

Docket No. 78414-A/JPW/GJG/ML

IN THE UNITED STATES PATENT AND TRADEMARK OFFICERECEIVED  
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Applicants : Nurit Livnah et al.

Serial No. : 10/764,288

Filing Date : January 23, 2004

For : PROTEIN KINASE INHIBITORS COMPRISING ATP  
MIMETICS CONJUGATED TO PEPTIDES OR  
PEPTIDOMIMETICS

JAN 23 2008

1185 Avenue of the Americas  
New York, New York 10036  
January 23, 2008BY FACSIMILE - 571-273-8300Mail Stop Petition  
Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450  
Attn: Ms. Karen Creasy  
Office of Petitions

SIR:

PETITION FOR WITHDRAWAL FROM  
ISSUE UNDER 37 C.F.R. §1.313(c)(2)

Applicant hereby petitions that U.S. Serial No. 10/764,288 be withdrawn from issue under 37 C.F.R. §1.313(c)(2) for consideration of a Request For Continued Examination ("RCE") in compliance with 37 C.F.R. §1.114 which applicant is filing concurrently herewith to list the divisional of the subject application on a Form PTO-1449. The RCE is attached hereto as Exhibit A and is accompanied by a Second Supplemental Information Disclosure Statement.

The required fee set forth in 37 C.F.R. §1.17(h) is \$130.00, and a check for \$940.00 including this amount is enclosed.

The PTO did not receive the following  
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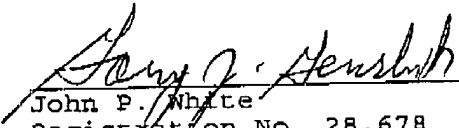
Applicants: Nurit Livnah, et al.  
Serial No. : 10/764,288  
Filed : January 23, 2004  
Page 2

Pursuant to 37 C.F.R. §1.313(c)(2), this Petition is being filed after payment of the issue fee on December 19, 2007 to request consideration of the accompanying RCE and Information Disclosure Statement.

No fee, other than the \$130.00 Petition fee and the \$810.00 RCE filing fee, is deemed necessary in connection with the filing of this Petition. However, if any additional fee is required, authorization is hereby given to charge the amount of such fee to Deposit Account No. 03-3125.

If a telephone interview would be of assistance in advancing prosecution of the subject application, applicant's undersigned attorneys invite the Examiner to telephone them at the number provided below.

Respectfully submitted,

  
John P. White  
Registration No. 28,678  
Gary J. Gershik  
Registration No. 39,992  
Attorneys for Applicants  
Cooper & Dunham LLP  
1185 Avenue of the Americas  
New York, New York 10036  
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**EXHIBIT A**  
**Applicants: Nurit Livnah et al.**  
**Application No.: 10/764,288**  
**Filed: January 23, 2004**

Jan-23-08 03:07pm From-

+212-391-0631

T-700 P.001

F-825

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ATTORNEYS AT LAW

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**MESSAGE:**

PETITION FOR WITHDRAWAL FROM ISSUE UNDER 37 C.F.R. §1.313 (C) (2) for Nurit Livnah et al., PROTEIN KINASE INHIBITORS COMPRISING ATP MIMETICS CONJUGATED TO PEPTIDES OR PEPTIDOMIMETICS, Serial No. 10/764,288, filed January 23, 2004, including RCE Transmittal (in duplicate) (Exhibit A), a Supplemental Information Disclosure Statement, a PTO-1449 (Exhibit B), and copy of the reference (Exhibit 1).

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Docket No. 2609/78414-A/JFW/GJG/ML

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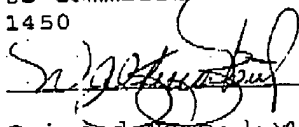
Applicants : Nurit Livnah et al.  
Serial No. : 10/764,288 Group Art Unit: 1654  
Filed : January 23, 2004 Examiner: A. Gupta  
For : PROTEIN KINASE INHIBITORS COMPRISING ATP  
MIMETICS COJUGATED TO PEPTIDES OR  
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January 23, 2008

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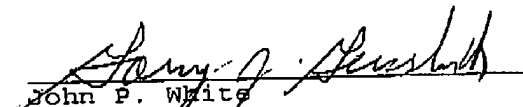
FACSIMILE CERTIFICATE OF TRANSMISSION  
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Date of Facsimile: January 23, 2008. I hereby certify that this paper is being transmitted to the Assistant Commissioner for Patents on the date indicated above by facsimile and is addressed to Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450



Printed Name: WALKIRIA BENITEZ

Respectfully submitted,



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Docket No. 78414-A/JPW/GJG/ML

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Commissioner for Patents  
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Alexandria, VA 22313-1450

SIR:

Second Supplemental Information Disclosure Statement

In accordance with their duty of disclosure under 37 C.F.R. §1.56, applicants would like to direct the Examiner's attention to the following reference, which is listed on Form PTO-1449 (Exhibit B).

This Second Supplemental Information Disclosure Statement is being submitted prior to the mailing of the first Office Action after the filing of a Request for Continued Examination under C.F.R. §1.114. Pursuant to 37 C.F.R. §1.97(b), this Information Disclosure Statement should be entered and considered.

1. U.S. Patent Application Serial No. 11/862,542, filed September 27, 2007 (Livnah et al.; divisional of the subject application) (Exhibit 1).

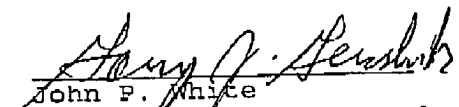
Applicants: Nurit Livnah, et al.  
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Page 2

Applicants request that the Examiner review the reference and make it of record in the subject application.

If a telephone interview would be of assistance in advancing prosecution of the subject application, applicants' undersigned attorney invites the Examiner to telephone him at the number provided below.

No fee is deemed necessary in connection with the filing of this Second Supplemental Information Disclosure Statement. However, if any other fee is required, authorization is hereby given to charge the amount of any such fee to Deposit Account No. 03-3125.

Respectfully submitted,

  
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**EXHIBIT B**

**Applicants: Nurit Livnah et al.**

**Application No.: 10/764,288**

**Filed: January 23, 2004**





# EXHIBIT 1

Applicants: Nurit Livnah et al.

Application No.: 10/764,288

Filed: January 23, 2004

**PROTEIN KINASE INHIBITORS COMPRISING ATP MIMETICS  
CONJUGATED TO PEPTIDES OR PEPTIDOMIMETICS**

**CROSS REFERENCE TO RELATED APPLICATIONS**

5           This application is a division of application no. 10/764,288 filed January 23, 2004, which is continuation of International application PCT/IL02/00618 filed July 25, 2002, the entire content of each of which is expressly incorporated herein by reference thereto.

**FIELD OF THE INVENTION**

10           The present invention relates to ATP mimetics, particularly isoquinoline derivatives, conjugated to a peptide or peptidomimetic, pharmaceutical compositions containing the isoquinoline derivatives and conjugates, their use as inhibitors of protein kinase, as well as to processes for the preparation and use of such molecules.

**BACKGROUND OF THE INVENTION**

15           Protein kinases are involved in signal transduction pathways linking growth factors, hormones and other cell regulation molecules to cell growth, survival and metabolism under both normal and pathological conditions. The superfamily of protein kinases includes protein kinase A and protein kinase C, as well as the more recently discovered protein kinase B (PKB).  
20           PKB is a newly recognized anti-apoptotic protein kinase whose activity is strongly elevated in human malignancies. PKB was originally discovered as a viral oncogene v-Akt in rat T-cell leukemia. It was later established that v-Akt is the oncogenic version of a cellular enzyme PKB/c-Akt, in which a truncated viral group specific antigen, gag, is fused in frame to the full length Akt-1 and is membrane bound whereas PKB/c-Akt is cytoplasmic. Sequencing of Akt  
25           revealed a high degree of homology to PKA (~75%) and PKC isozymes (~50%), a fact which led to its renaming as PKB.

          PKB activation involves phosphorylation of two amino acid residues, Ser473 and Thr308. The enzyme is activated by the second messenger PIP3 produced by PI-3-kinase. PIP3 binds to the pleckstrin homology (PH) domains of PKB, recruits it to the membrane where it is  
30           phosphorylated and converted to its activated form. Since PKB activation is PI-3-kinase dependent, the persistent activation of certain protein tyrosine kinases, such as IGF-1 receptor, EGF receptor, PDGF receptor, pp60c-Src, and the like, leads to the persistent activation of PKB.

which is indeed encountered in many tumors. Deletions in the gene coding for the tumor suppressor PTEN also induce the persistent activation of PKB/cAkt since it is the negative regulator of this enzyme. Also, PKB is overexpressed in 15% of ovarian cancers, 12% of pancreatic cancers and 3% of breast cancers, and was shown to produce a survival signal that protects cells from apoptosis thus contributing to resistance to chemotherapy.

PKB has emerged as a crucial regulator of widely divergent cellular processes including apoptosis, proliferation, differentiation and metabolism. Disruption of normal PKB/Akt signaling has now been documented as a frequent occurrence in several human cancers and the enzyme appears to play an important role in their progression (Nicholson and Anderson, Cellular Signalling 14, 381, 2002). Therefore PKB is, in principle, an attractive drug target for the treatment of cancer. Ideally, a drug that inhibits PKB should cause both cell cycle arrest and promote apoptosis. Such activity would result in increased cell death of tumor tissue where PKB is elevated, and in decreased resistance to chemotherapy agents.

These molecular properties of PKB and its central role in tumorigenesis, implies that this protein kinase may be an attractive target for novel anti-cancer agents. To date no specific inhibitors of PKB are known in the art, nor are any of the disclosed inhibitors of protein kinases A and C known to act on PKB.

Hidaka H. et al. (Biochemistry, 32, 5036, 1984) describe a class of isoquinolinesulfonamides having inhibitory activity towards cyclic nucleotide dependent protein kinases (PKA and PKG) and protein kinases C (PKC). The same class of compounds is claimed in EP 061673, which discloses said compounds as having cardiovascular activity. Additional derivatives of isoquinolinesulfonyl were disclosed by Hidaka in EP 109023, US 4456757, US 4525589, and US 4560755.

Antitumor activity has been suggested for some of these isoquinolinesulfonamides. Martell R.E. et al. (Biochem. Pharm., 37, 635, 1988) found effects of two isoquinolinesulfonamides, namely 1-(5-isoquinolinsulfonyl)-2-methylpiperazine (H-7) and N-[2-guanidinoethyl]-5-isoquinolinesulfonamide (HA-1004), which have a certain selectivity for PKC and cyclic nucleotide dependent protein kinases, respectively, on calcitriol-induced cell differentiation. Further, Nishikawa M. et al., Life Sci., 39, 1101, 1986), demonstrate that the same compound H-7 inhibits cell differentiation induced by phorbol diester.

International PCT application WO 93/13072 discloses 5-isoquinolinesulfonamide derivatives as protein kinase inhibiting agents wherein the claimed compounds all contain two sulfonyl moieties.

Other classes of compounds known in the prior art (EP-A-397060, DE-A-3914764 and EP-A-384349) showed the capacity of inhibiting protein kinases, however, said compounds have a chemical structure which is totally different from that of the compounds of the present invention. In addition, international PCT application WO 98/53050 discloses short peptides derived from the HJ loop of a serine/threonine kinase which modulate the activity of serine/threonine kinases.

The minimal consensus sequence for efficient phosphorylation by PKB was found by Alessi et al. (Fed. Eur. Biochem. Soc., 399, 333, 1996). This is a 7-mer motif with the most active peptide substrate having the sequence Arg—Pro—Arg—Thr—Ser—Ser—Phe (SEQ ID NO: 1). International application WO 97/22360 discloses certain PKB substrate peptides having 7-amino acids length, useful as substrate for measuring PKB activity.

Obata et al. (J. Biol. Chem., 17, 36108, 2000) described the use of an oriented peptide library approach to determine optimal amino acid sequence of the PKB substrate. All the substrates identified contained the known motif having the sequence Arg—Xaa—Arg—Xaa—Xaa—Ser/Thr (SEQ ID NO: 2).

Ricouart et al. (J. Med. Chem. 1991, 34, 73-78), described conjugates of isoquinolinesulfonamides and peptides for the inhibition of PKA. Loog et al. (Bioorganic and Medicinal Chemistry Letters 1999, 9, 1447-1452), described a chimera with adenosine and peptides for the inhibition of PKA and PKC. The inhibition obtained with the disclosed compound is very poor. Schlitzer et al. (Bioorganic and Medicinal Chemistry, 2000, 1991-2006) deal with a small molecule linked to non-peptidic long chain moieties that are supposed to replace the peptide part of the substrate. The disclosed compounds show poor inhibitory activity.

Parang et al. (Nature Structural Biology 8, 37, 2001), describe peptide-ATP bisubstrate analogs of a protein kinase A inhibitor, wherein ATP is linked to a protein kinase peptide substrate. Nevertheless, this approach has a limitation of suboptimal pharmacokinetic properties. WO 01/70770 discloses bisubstrate inhibitors for the insulin receptor tyrosine kinase, and a specific potent and selective inhibitor comprising an ATP-gamma-S linked to a peptide substrate analog via a two-carbon spacer.

Numerous disclosures in the background art and in co-pending International Patent Application WO 01/91754 by one of the present inventors relate to specific isoquinoline derivatives, which are PKB inhibitors. The present invention is directed to novel isoquinoline derivatives and more specifically isoquinoline conjugates, and excludes all known compounds previously claimed for their capacity to inhibit PKB.

The present invention overcomes the limitations of known inhibitors by providing ATP surrogates and peptidomimetics with protein target specificity.

### SUMMARY OF THE INVENTION

It is an object of the present invention to provide specific inhibitors of protein kinases for medical, therapeutic and drug design purposes. It is yet another object of the present invention to provide such molecules, which are selective inhibitors of protein kinase B.

One aspect of the present invention involves the preparation of novel compounds which inhibit the activity of protein kinases. It has now been found that certain novel derivatives of isoquinolinesulfonamides, which are protein kinase inhibiting agents, when conjugated to a peptide or peptidomimetic, unexpectedly proved to be active towards a specific type of protein kinase, namely protein kinase B.

The present invention provides small molecules having high affinity to the ATP binding site of PKB, which are conjugated to a peptide or peptidomimetic moiety which mimics the substrate of PKB. These compounds are referred to herein as "chimeric" compounds. The chimeric compounds according to the present invention preferably serve as PKB inhibitors with improved activity and selectivity.

Another aspect of the present invention is directed to pharmaceutical compositions comprising as an active ingredient novel inhibitors of protein kinase and to methods for the preparation and use of pharmaceutical compositions comprising these novel inhibitors of protein kinases.

Another aspect of the present invention is directed to the use of pharmaceutical compositions comprising these protein kinase inhibitors for production of medicaments useful for the treatment or diagnosis of diseases and disorders. The present invention discloses methods of treatment of disorders involving protein kinase, including but not limited to cancers,

cardiovascular pathologies, hemorrhagic shock, obesity, inflammatory diseases, diseases of the central nervous system, and autoimmune diseases.

The present invention further provides methods for modulating the activity of protein kinases in a subject, comprising administering a therapeutically effective amount of a protein kinase inhibitor.

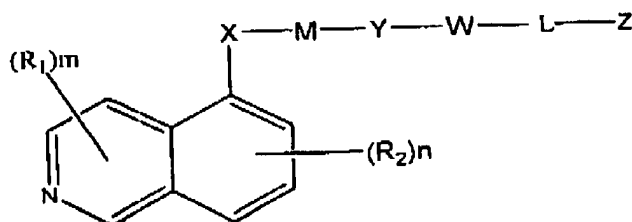
Further aspects of the present invention are directed to methods for the diagnosis of diseases including in-vitro diagnosis using the compounds of the present invention, and in-vivo diagnosis comprising administering a pharmaceutical composition comprising a diagnostically useful amount of a protein kinase inhibitor prepared according to the principles of the present invention.

It is yet another object of the present invention to provide protein kinase inhibitors comprising peptidomimetic compounds having improved stability and cell permeability properties. Non limiting examples of such compounds include N-alkylation of selected peptide residues, side-chain modifications of selected peptide residues, non-natural amino acids, use of carbamate, urea, sulfonamide and hydrazine for peptide bond replacement, and incorporation of non-peptide moieties including but not limited to piperidine, piperazine and pyrrolidine, through a peptide or non-peptide bond. These peptidomimetic compounds may be used according to the present invention as the peptide substrate part of chimeric compounds. In addition these peptidomimetic compounds may be used as protein kinase inhibitors per se.

Preferred embodiments according to the present invention comprise a chimeric compound comprising both an ATP mimetic moiety and a peptidic substrate mimetic moiety connected by a spacer.

The ATP mimetic core includes but is not limited to dansyls, isoquinolines, quinolines and naphthalenes. The spacer is of varied lengths and conformations of any suitable chemistry including but not limited to amine, amide, thioether, oxyether, sulfonamide bond and the like. Non limiting examples for such spacers include sulfone amide derivatives, amino thiol derivatives and amino alcohol derivatives. The peptidic moiety comprises peptides or peptidomimetics. Such inhibitory peptides are designed based on any peptide which may serve as a PKB substrate.

More preferred embodiments of the present invention comprise a compound of Formula I:



Formula I

wherein:

- 5  $R_1$  and  $R_2$  are independently selected from the group consisting of hydrogen, a lower alkyl group, a lower alkoxy group, substituted or unsubstituted phenyl group, a lower alkyl substituted with at least one substituent selected from the group consisting of a phenyl group, a halogen, hydroxyl, thiol, nitro, cyano, or amino group;  
 $m$  and  $n$  are each independently 0-3;
- 10  $X$  is selected from the group consisting of  $SO_2-NH$ ,  $S$  and  $O$ ;  
 $M$  represents substituted or unsubstituted alkylene of 1-4 carbon atoms;  
 $Y$  is selected from the group consisting of amide, amine, urea, carbamate, hydrazine or sulfonamide;
- 15  $W$  is absent or is selected from the group consisting of substituted or unsubstituted alkylene, aliphatic, aromatic or heterocyclic moiety, of 1-18 carbon atoms;  
 $L$  is absent or is selected from the group consisting of amide, amine, urea, carbamate, hydrazine or sulfonamide; and  
 $Z$  is a peptide or peptidomimetic moiety of 4-12 residues in length capable of binding to the substrate site of PKB.
- 20 Preferably,  $R_1$  and  $R_2$  are independently selected from the group consisting of methyl, ethyl, ethoxy and dimethylamine;  
 $m$  and  $n$  are each 1;  
 $X$  is selected from the group consisting of  $SO_2-NH$  and  $S$ ;  
 $M$  represents substituted or unsubstituted alkylene of 2 carbon atoms;
- 25  $Y$  is selected from the group consisting of amide and amine;  
 $W$  is absent or is selected from the group consisting of substituted or unsubstituted alkylene, aliphatic, aromatic or heterocyclic moiety, of 1-5 carbon atoms;  
 $L$  is absent or is selected from the group consisting of amide and amine; and



Z is a peptide or peptidomimetic moiety of 6-10 residues in length capable of binding to the substrate site of PKB.

According to certain currently more preferred embodiments the peptide substrate mimetic designated as Z in Formula I, comprises a sequence of seven residues that are referred to herein as AA<sub>1</sub> through AA<sub>7</sub>. Each AA is selected from the group consisting of an amino acid, an amino acid analog, or an aliphatic, aromatic or heterocyclic moiety, incorporated into the sequence to create a peptidomimetic moiety with improved pharmacological properties.

AA<sub>1</sub> and AA<sub>3</sub> are independently selected from the group consisting of: arginine or arginine analog; lysine or lysine analog; ornithine or ornithine analog; or an aliphatic, aromatic, or heterocyclic moiety bearing a group positively charged at physiological pH, such as an amine, guanidine or amidine, homoarginine, arginino.

AA<sub>2</sub> is selected from the group consisting of proline, proline analog or an aliphatic, aromatic or heterocyclic moiety, hydroxyproline, nipecotic acid, alanine, aminobutyric acid.

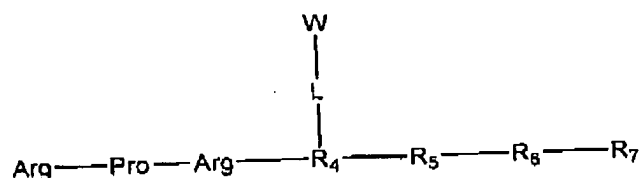
AA<sub>4</sub>, AA<sub>5</sub>, AA<sub>6</sub> are each independently selected from the group consisting of: diaminopropionic acid, diaminobutyric acid, ornithine, GlyNH<sub>2</sub>, Tyr or Tyr analog, Thr or Thr analog; Ser or Ser analog; Ala or Ala analog; Glu or Glu analog, Gly or Gly analog; an aliphatic, aromatic or heterocyclic residue bearing alkyl, benzyl, hydroxy, phenoxy alkoxy, sulfone, sulfoxide, phosphonate, phosphonate ester, amide or carbamoyl functionality, amino butyric acid, citrulline, serinol, phosphotyrosine and phosphotyrosine dimethyl ester.

AA<sub>7</sub> is selected from the group consisting of: Phe or Phe analog; Trp or an analog, Tyr or an analog, Leu or an analog, homoleucine or an analog, Ile or an analog; aromatic moiety esters or aromatic substitutions of an amino acid; an aromatic, heterocyclic or branched aliphatic moiety; homophenylalanine, homoleucine, glutamic benzyl ester, naphthylalanine.

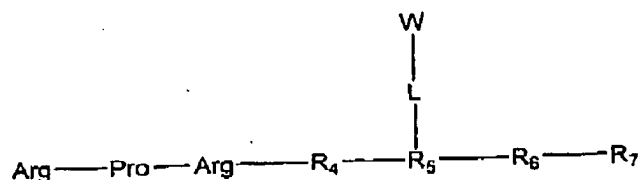
Due to the peptidomimetic nature of the preferred embodiments according to the invention, the bonds between AAs may be selected from the group consisting of: an amide, urea, carbamate, hydrazine or sulfonamide bond. In the currently more preferred embodiments the bonds between the AAs are all amide bonds unless explicitly stated otherwise.

Additional preferred embodiments of the present invention comprise a compound of Formulae IIa - IIc:

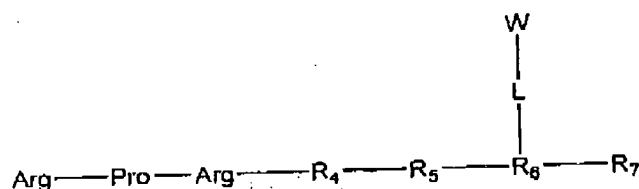
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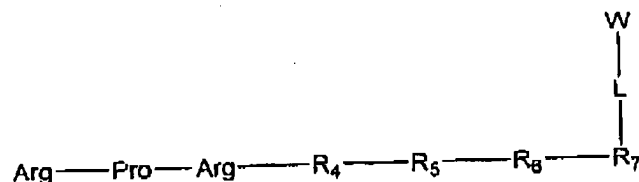
Formula IIa



Formula IIb



Formula IIc



Formula IId

5

wherein:

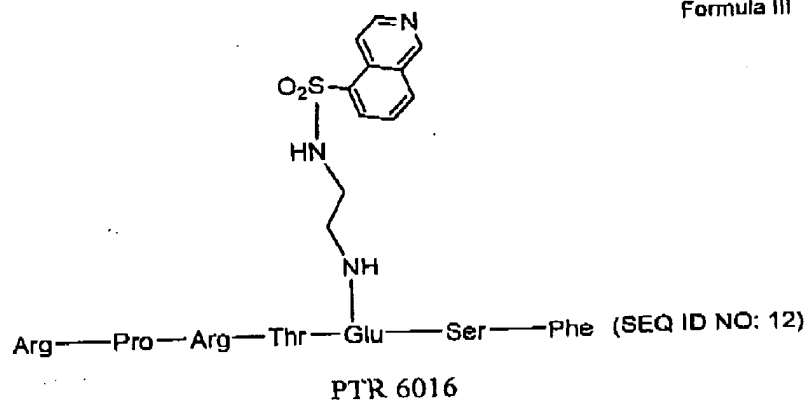
10  $R_4$ ,  $R_5$ , and  $R_6$  are each independently selected from the group consisting of threonine, serine, glutamic acid allyl ester, homocitrulline, lysine, methionine, norleucine, ornithine, arginine, glycine, diaminopropionic acid, diaminobutyric acid, GlyNH<sub>2</sub>, and alanine; or are an N<sup>α</sup>-ω-functionalized derivative of an amino acid selected from the group of glycine, alanine and tyrosine.

$R_7$  is selected from the group consisting of phenylalanine, homoleucine, norleucine, glutamic acid allyl ester;

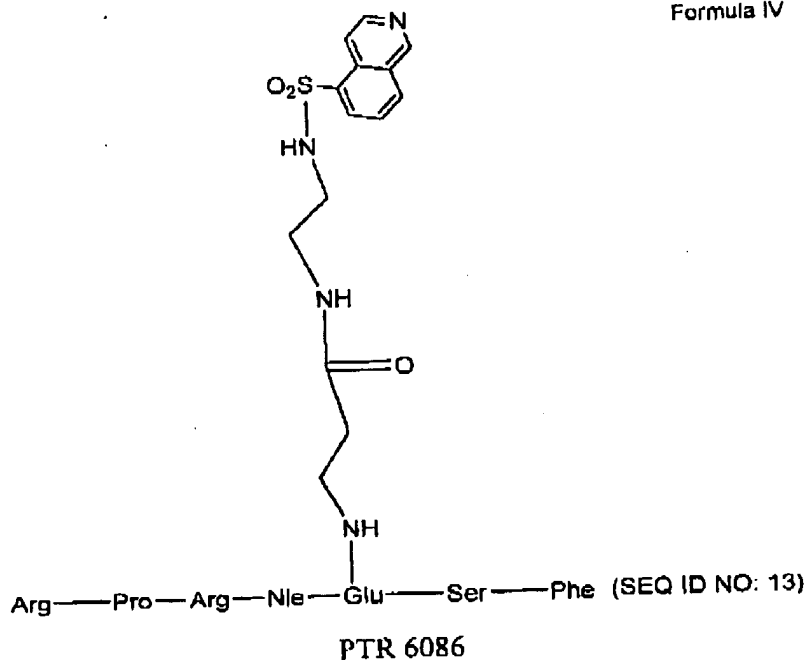
W is absent or is N-(8-sulfonamide-5-isoquinoline) ethylenediamine; and  
 L may be absent or is selected from the group consisting of glycine,  $\beta$ -alanine, phenylalanine,  
 aminobutyric acid and aminopentanoic acid.  
 Preferably, W is connected to  $R_5$  as described in Formula IIb.

5 Currently most preferred embodiments of the present invention are the chimeric  
 compounds of Formulae III-VII:

Formula III

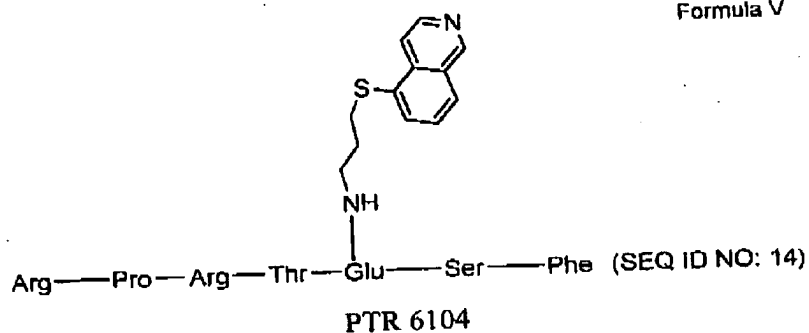


Formula IV

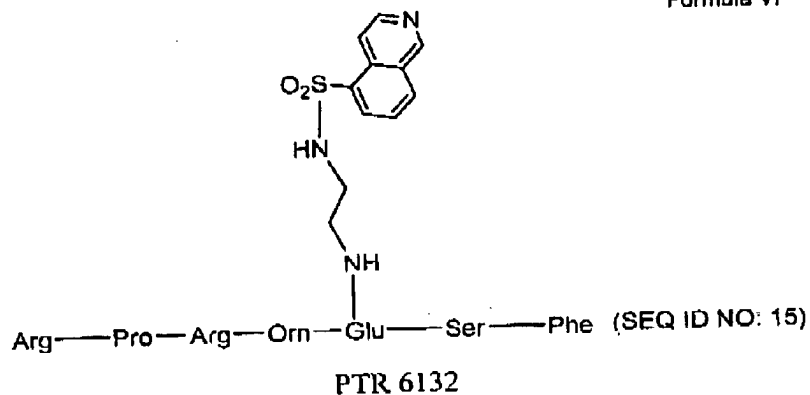


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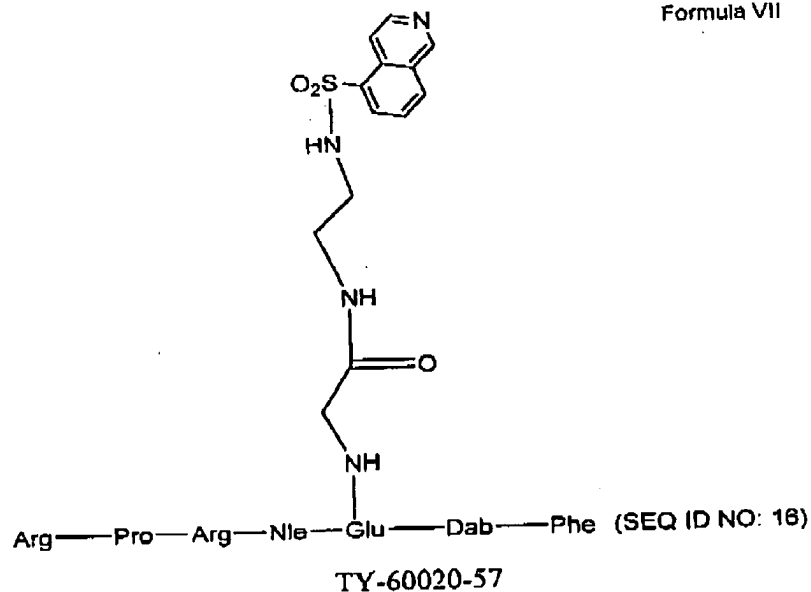
Formula V



Formula VI



Formula VII



Currently most preferred are compounds having a sequence selected from:

Arg—Pro—Arg—Thr—Glu—(bAla-5-mercaptoaminopropyl-isoquinoline)—Ser—Phe  
(SEQ ID NO: 3).

Arg—Pro—Arg—Thr—Glu—(5-mercaptoaminopropyl-isoquinoline)—Ser—Phe (SEQ  
5 ID NO: 4).

Arg—Pro—Arg—Orn—Glu—(5-aminoethylsulfonamide isoquinoline)—Ser—Phe (SEQ  
ID NO: 5).

Arg—Pro—Arg—Nva—Glu—(5-mercaptoaminopropyl-isoquinoline)—Ser—Phe (SEQ  
ID NO: 6).

Arg—Pro—Arg—Nle—Glu—(5-mercaptoaminopropyl-isoquinoline)—Ser—Phe (SEQ  
10 ID NO: 7).

Arg—Pro—Arg—Orn—Glu—(Gly-5-aminoethylsulfonamide)—Dab—Hol (SEQ ID  
NO: 8).

Arg—Pro—Arg—Nle—Glu—(Gly-5-aminoethylsulfonamide)—Dab—Phe (SEQ ID  
15 NO: 9).

Arg—Pro—Arg—Nle—Glu—(Gly-5-aminoethylsulfonamide)—Dab—Hol (SEQ ID  
NO: 10).

20 Essentially all of the uses known or envisioned in the prior art for protein kinase inhibitors, can be accomplished with the molecules of the present invention. These uses include therapy and diagnostic techniques.

By way of exemplification, the compounds disclosed in the present invention were selected for inhibition of Protein kinase B. Using the preparations and methods disclosed herein it is possible to obtain compounds that inhibit the activity of other types of protein kinases.

25

### BRIEF DESCRIPTION OF THE DRAWINGS

**Figure 1.** The synergistic effect of chimeric compound PTR 6016 composed of an ATP mimic and a peptide substrate.

**Figure 2.** Inhibition activity curve for AR-60003-52 as described in example 6.

30 **Figure 3.** Describes PKB and PKA inhibition curves (IC<sub>50</sub>) of the active and selective peptide denoted 60018-16.

## DETAILED DESCRIPTION OF THE INVENTION

It is now disclosed that chimeric compounds according to the present invention are inhibitors of protein kinases. It is known that small molecules that share certain structural motifs may serve as "ATP mimetic" inhibitors. It has now been discovered that such moieties may be  
5 part of chimeric compounds comprising also a substrate mimetic moiety, which are potent inhibitors of protein kinases, particularly PKB, according to the present invention.

The disclosed protein kinase inhibitors are chimeric molecules which exhibit enhanced affinity toward certain protein kinase subtypes. In principle, the present invention provides for  
10 the first time highly active (in the nano molar range) inhibitors of protein kinase B. The preferred molecules generally have a molecular weight of less than about 1100 daltons. These and further advantages over the background art will become apparent from the description of the currently preferred embodiments of the present invention.

Preferred compounds according to the present invention are composed of small molecules having high affinity to the ATP binding site of PKB, which are conjugated to a  
15 peptide or peptido-mimetic moiety that mimics the substrate of PKB. The chimeric compounds according to the present invention preferably serve as PKB inhibitors with improved activity and selectivity.

The utility of the compositions according to the invention can be established by means of various assays as are well known in the art. The preferred compounds of the present invention  
20 were found to be active in a panel of in-vitro assays, in inhibiting the activity of protein kinases and in induction of apoptosis in cancer cells.

Pharmaceutical compositions according to the present invention comprising pharmacologically active protein kinase inhibitors and a pharmaceutically acceptable carrier or diluent represent another embodiment of the invention, as do the methods for the treatment of a  
25 mammal in need thereof with a pharmaceutical composition comprising an effective amount of a protein kinase inhibitor according to the invention. Methods of treatment using the compositions of the invention are useful for therapy of cancers, diabetes, cardiovascular pathologies, hemorrhagic shock, obesity, inflammatory diseases, diseases of the central nervous system, and autoimmune diseases using such compositions.

30 The pharmaceutical compositions according to the present invention may be most preferably be used for prevention and treatment of malignancies selected from the group of

Breast Cancer (Perez-Tenorio and Stal, Br.J.cancer 2002 86, 540-45, Salh et al, Int. J.cancer 2002 98,148-54); Ovarian cancer (Liu et al, cancer res. 1998 15, 2973-7); Prostate cancer (Zin et al, Clin.cancer.res.2001 7,2475-9); Colon cancer (Semba at al, clin.cancer.res. 2002 8,1957-63); Melanoma and skin cancer (Walderman, Wecker and Diechmann, Melanoma res. 2002 12, 45-50); Lung cancer(Zin et al, Clin.cancer.res.2001 7,2475-9); and hepatocarcinoma (Fang et al, Eur. J. Biochem. 2001 268, 4513-9).

The pharmaceutical compositions according to the present invention advantageously comprise at least one protein kinase inhibitor. These pharmaceutical compositions may be administered by any suitable route of administration, including topically or systemically. Preferred modes of administration include but are not limited to parenteral routes such as intravenous and intramuscular injections, as well as via nasal or oral ingestion.

As is known to those skilled in the art the pharmaceutical compositions may be administered alone own or in conjunction with additional treatments for the conditions to be treated.

15

#### Terminology and definitions

In the specification and in the claims the term "protein kinase" refers to a member of an enzyme superfamily which functions to phosphorylate one or more protein as described above.

As used herein and in the claims, the term "inhibitor" is interchangeably used to denote "antagonist" these terms define compositions which have the capability of decreasing certain enzyme activity or competing with the activity or function of a substrate of said enzyme.

As used herein and in the claims the term "chimeric compound" or "chimeric molecule" denotes an ATP mimic moiety conjugated to a PKB substrate mimetic part. Examples for such chimeric compounds or conjugates are small molecules (and more specific isoquinoline derivatives) that mimic the ATP molecule of the PKB, conjugated to a peptide or a peptidomimetic moiety which is a PKB substrate mimetic. These molecules may preferably serve as PKB inhibitors with improved activity and selectivity.

As used herein "peptide" indicates a sequence of amino acids linked by peptide bonds. The peptide analogs of this invention comprise a sequence of 3 to 15 amino acid residues, preferably 4 to 12 residues, more preferably 5 to 10 amino acids, each residue being characterized by having an amino and a carboxy terminus.

The term "peptidomimetic" means that a peptide according to the invention is modified in such a way that it includes at least one non-coded residue or non-peptidic bond. Such modifications include, e.g., alkylation and more specific methylation of one or more residues, insertion of or replacement of natural amino acid by non-natural amino acids, replacement of an amide bond with other covalent bond. A peptidomimetic according to the present invention may optionally comprises at least one bond which is an amide-replacement bond such as urea bond, carbamate bond, sulfonamide bond, hydrazine bond, or any other covalent bond. The design of appropriate "peptidomimetic" may be computer assisted.

The term "peptide analog" indicates molecule which has the amino acid sequence according to the invention except for one or more amino acid changes or one or more modification/replacement of an amide bond.

In the specification and in the claims the term "therapeutically effective amount" refers to the amount of protein kinase inhibitor or composition comprising same to administer to a host to achieve the desired results for the indications described herein, such as but not limited of cancers, diabetes, cardiovascular pathologies, hemorrhagic shock, obesity, inflammatory diseases, diseases of the central nervous system, and autoimmune diseases.

Certain abbreviations are used herein to describe this invention and the manner of making and using it. For instance, ATP refers to adenosine three phosphate, BSA refers to bovine serum albumin, BTC refers to bis-(trichloromethyl)carbonate or triphosgene, DCM refers to dichloromethane, DIEA refers to diisopropyl-ethyl amine, DMF refers to dimethyl formamide, EDT refers to ethanedithiol, EDTA refers to ethylene diamine tetra acetate, ELISA refers to enzyme linked immuno sorbent assay, EGF refers to epithelial growth factor, FACS refers to fluorescence assisted cell sorter, HA refers to hemagglutinin, HBTU refers to 1-hydroxybenzotriazolyltetramethyl-uronium, HEPES refers to 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid, HOBt refers to 1-hydroxybenzotriazole, HRP refers to horse raddish peroxidase, IGF refers to insulin growth factor, MOPS refers to 4-morpholinepropanesulfonic acid, MPS refers to multiple parallel synthesis, NMP refers to N-methyl formamide, OPD refers to o-Phenylenediamine, PBS refers to phosphate buffer saline, PKA refers to protein kinase A, PKB refers to protein kinase B, PKC refers to protein kinase C, rpm refers to rounds per minute, SAR refers to structure-activity relationship, THF refers to tetrahydrofuran, TIS refers to tri-isopropyl-silane, TFA refers to trifluoric acetic acid.



The amino acids used in this invention are those which are available commercially or are available by routine synthetic methods. Certain residues may require special methods for incorporation into the peptide, and either sequential, divergent or convergent synthetic approaches to the peptide sequence are useful in this invention. Natural coded amino acids and their derivatives are represented by three-letter codes according to IUPAC conventions. When there is no indication, the L isomer was used. The D isomers are indicated by "D" before the residue abbreviation.

Conservative substitution of amino acids as known to those skilled in the art are within the scope of the present invention. Conservative amino acid substitutions includes replacement of one amino acid with another having the same type of functional group or side chain e.g. aliphatic, aromatic, positively charged, negatively charged.

List of non limiting examples of non-coded amino acids which where used in the present invention: Abu refers to 2-aminobutyric acid, Ape5 refers to aminopentanoic acid, ArgOl refers to argininol, bAla refers to  $\beta$ -Alanine, Bpa refers to 4-Benzoylphenylalanine, Bip refers to Beta-(4-biphenyl)-alanine, Dab refers to diaminobutyric acid, Dap refers to Diaminopropionic acid, Dim refers to Dimethoxyphenylalanine, Dpr refers to Diaminopropionic acid, Hol refers to homoleucine, HPh refers to Homophenylalanine, Gaba refers to gamma aminobutyric acid, GlyNH<sub>2</sub> refers to Aminoglycine, Nie refers to Norleucine, Nva refers to Norvaline, Orn refers to Ornithine, PheCarboxy refers to para carboxy Phenylalanine, PheCl refers to para chloro Phenylalanine, PheF refers to para fluoro Phenylalanine, PheMe refers to para methyl Phenylalanine, PheNH<sub>2</sub> refers to para amino Phenylalanine, PheNO<sub>2</sub> refers to para nitro Phenylalanine, Phg refers to Phenylglycine, Thi refers to Thienylalanine.

#### Pharmacology

The compounds of the present invention can be administered to a subject in a number of ways, which are well known in the art. Hereinafter, the term "subject" refers to the human or lower animal to whom compounds of the present invention are administered.

The novel pharmaceutical compositions of the present invention contain in addition to the active ingredient conventional pharmaceutically acceptable carriers, diluents and the like. Solid compositions for oral administration such as tablets, pills, capsules or the like may be prepared by mixing the active ingredient with conventional, pharmaceutically acceptable ingredients such

as corn starch, lactose, sucrose, sorbitol, talc, stearic acid, magnesium stearate, dicalcium phosphate and gums with pharmaceutically acceptable diluents. The tablets or pills can be coated or otherwise compounded with pharmaceutically acceptable materials known in the art to provide a dosage form affording prolonged action or sustained release. Other solid compositions  
5 can be prepared as suppositories, for rectal administration.

Liquid forms may be prepared for oral administration or for injection, the term including subcutaneous, transdermal, intravenous, intrathecal, and other parenteral routes of administration. The liquid compositions include aqueous solutions, with or without organic cosolvents, aqueous or oil suspensions, emulsions with edible oils, as well as other micellar  
10 dispersions and similar pharmaceutical vehicles. In addition, the compositions of the present invention may be formed as aerosols, for intranasal and like administration. More preferred formulations include sustained release or depot formulations, which may provide a steady state pharmacokinetic profile.

However, it is evident to the man skilled in the art that dosages would be determined by  
15 the attending physician, according to the disease to be treated, method of administration, patient's age, weight, contraindications and the like.

All the compounds defined above are effective as inhibitors of protein kinase and can be used as active ingredients of pharmaceutical compositions for treatment of one, or simultaneously several, symptoms of the disorders defined above.

20 The compounds of the present invention are administered for the above-defined purposes in conventional pharmaceutical forms, with the required solvents, diluents, excipients, etc. to produce a physiologically acceptable formulation. They can be administered by any of the conventional routes of administration.

It will be appreciated that the most appropriate administration of the pharmaceutical  
25 compositions of the present invention will depend on the type of disorder or disease being treated.

#### Chemistry:

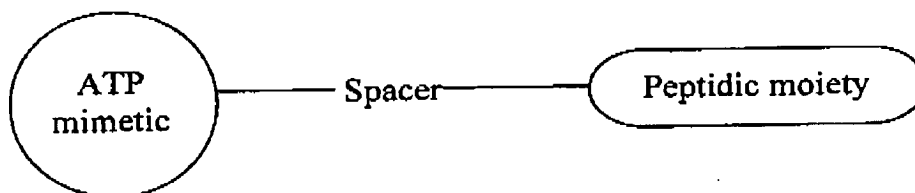
Some of the preferred compounds of the present invention may conveniently be prepared  
30 using solution phase synthesis methods. Other methods known in the art to prepare isoquinoline compounds like those of the present invention, can be used and are comprised in the scope of the

present invention. Preferred peptides according to the present invention may be synthesized using any method known in the art, including peptidomimetic methodologies. These methods include solid phase as well as solution phase synthesis methods. The conjugation of the peptidic and small molecule moieties may be performed using any methods known in the art, either by solid phase or solution phase chemistry. Non-limiting examples for these methods are described hereby.

By way of exemplification of the principles of the present invention, a search for inhibitory PKB chimeric compounds focused on SAR studies of certain molecules, as exemplified hereinbelow.

#### Preferred embodiments

Protein kinases have more than one active site, they possess a catalytic site for ATP and a substrate-binding site. Preferred compounds according to the present invention can bind both sites at the same time and may have a synergistic effect that will give it unique potency and selectivity properties. These preferred compounds are chimeric molecules which are designed to include an ATP-mimetic molecule, connected via various spacers to a substrate-mimetic portion. Preferred embodiments according to the present invention comprise a chimeric compound comprising both an ATP mimetic moiety and a peptidic substrate mimetic moiety connected by a spacer, as described in the following scheme I:



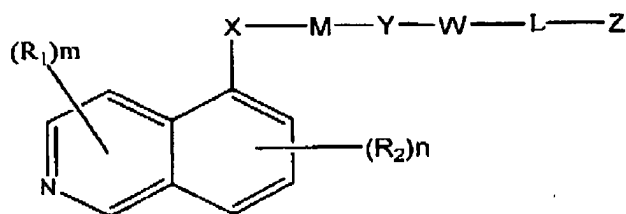
Scheme I

The ATP mimetic core includes but is not limited to dansyls, isoquinolines, quinolines and naphthalenes. The spacer is of varied lengths and conformations of any suitable chemistry including but not limited to amine, amide, thioether, oxyether, sulfonamide bond and the like.

Non-limiting examples for such spacers include sulfone amide derivatives, amino thiol derivatives and amino alcohol derivatives. The peptidic moiety comprises peptides and

peptidomimetics. Such inhibitory peptides may be designed based on any peptide which may serve as PKB substrate.

Additional more preferred embodiment of the present invention comprises a compound of Formula I:



Formula I

wherein:

$R_1$  and  $R_2$  are independently selected from the group consisting of hydrogen, a lower alkyl group, a lower alkoxy group, substituted or unsubstituted phenyl group, a lower alkyl substituted with at least one substituent selected from the group consisting of a phenyl group, a halogen, hydroxyl, thiol, nitro, cyano, or amino group;

$m$  and  $n$  are each independently 0-3;

$X$  is selected from the group consisting of  $SO_2-NH$ ,  $S$  and  $O$ ;

$M$  represents substituted or unsubstituted alkylene of 1-4 carbon atoms;

$Y$  is selected from the group consisting of amide, amine, urea, carbamate, hydrazine or sulfonamide;

$W$  is absent or is selected from the group consisting of substituted or unsubstituted alkylene, aliphatic, aromatic or heterocyclic moiety, of 1-18 carbon atoms;

$L$  is absent or is selected from the group consisting of amide, amine, urea, carbamate, hydrazine or sulfonamide; and

$Z$  is a peptide or peptidomimetic moiety of 4-12 residues in length capable of binding to the substrate site of PKB.

Preferably,  $R_1$  and  $R_2$  are independently selected from the group consisting of methyl, ethyl, ethoxy and dimethylamine;

$m$  and  $n$  are each 1;

$X$  is selected from the group consisting of  $SO_2-NH$  and  $S$ ;

M represents substituted or unsubstituted alkylene of 2 carbon atoms;

Y is selected from the group consisting of amide and amine;

W is absent or is selected from the group consisting of substituted or unsubstituted alkylene, aliphatic, aromatic or heterocyclic moiety, of 1-5 carbon atoms;

5 L is absent or is selected from the group consisting of amide and amine; and

Z is a peptide or peptidomimetic moiety of 6-10 residues in length capable of binding to the substrate site of PKB.

Preferred peptide substrate and peptide substrate mimetics according to the present invention, forming part of the chimeric compounds are described in the following scheme:

10



According to this scheme AA is selected from the group consisting of an amino acid, an amino acid analog, or an aliphatic, aromatic or heterocyclic moiety, incorporated into the sequence to create a peptidomimetic moiety with improved pharmacological properties.

15 AA<sub>1</sub> and AA<sub>3</sub> are independently selected from the group consisting of: arginine or arginine analog; lysine or lysine analog; ornithine or ornithine analog; or an aliphatic, aromatic, or heterocyclic moiety bearing a group positively charged at physiological pH, such as an amine, guanidine or amidine, homoarginine, argininol.

20 AA<sub>2</sub> is selected from the group consisting of proline, proline analog or an aliphatic, aromatic or heterocyclic moiety, hydroxyproline, nipecotic acid, alanine, aminobutyric acid.

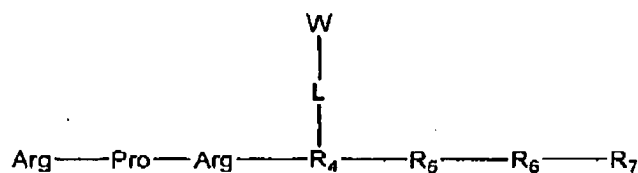
25 AA<sub>4</sub>, AA<sub>5</sub>, AA<sub>6</sub> are independently selected from the group consisting of: Diaminopropionic acid, diaminobutyric acid, Ornithine, GlyNH<sub>2</sub>, Tyr or Tyr analog, Thr or Thr analog; Ser or Ser analog; Ala or Ala analog; Glu or Glu analog, Gly or Gly analog; an aliphatic, aromatic or heterocyclic residue bearing alkyl, benzyl, hydroxy, phenoxy alkoxy, sulfone, sulfoxide, phosphonate, phosphonate ester, amide or carbamoyl functionality, amino butyric acid, citrulline, serinol, phosphotyrosine and phosphotyrosine dimethyl ester.

30 AA<sub>7</sub> is selected from the group consisting of: Phe or Phe analog; Trp, Tyr, Leu, HomoLeucine, Ile, and their analogs; aromatic moiety esters or aromatic substitutions of an amino acid; an aromatic, heterocyclic or branched aliphatic moiety; homophenylalanine, homoleucine, glutamic benzyl ester, naphthylalanine.

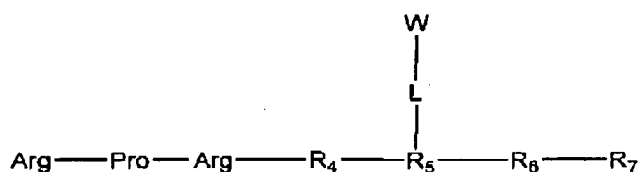
Due to the peptidomimetic nature of the preferred embodiments according to the invention, the bonds between AAs are not only peptide bonds but may be selected from the group consisting of: an amide, urea, carbamate, hydrazine or sulfonamide bond.

Additional preferred embodiments of the present invention comprise a compound of

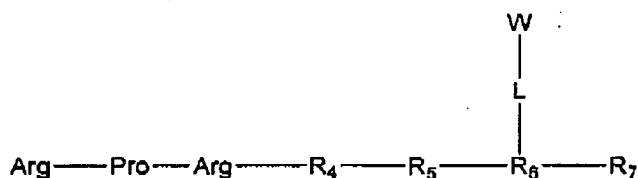
5 Formulae IIa - IId:



Formula IIa



Formula IIb



Formula IIc



Formula II d

10 wherein:

$R_4$ ,  $R_5$ , and  $R_6$  are each independently selected from the group consisting of threonine, serine, glutamic acid allyl ester, homocitrulline, lysine, methionine, norleucine, ornithine, arginine,

glycine, diaminopropionic acid, diaminobutyric acid, GlyNH<sub>2</sub>, and alanine; or are an N<sup>α</sup>-ω-functionalized derivative of an amino acid selected from the group of Glycine, Alanine and Tyrosine;

R<sub>7</sub> is selected from the group consisting of phenylalanine, homoleucine, norleucine, glutamic acid allyl ester;

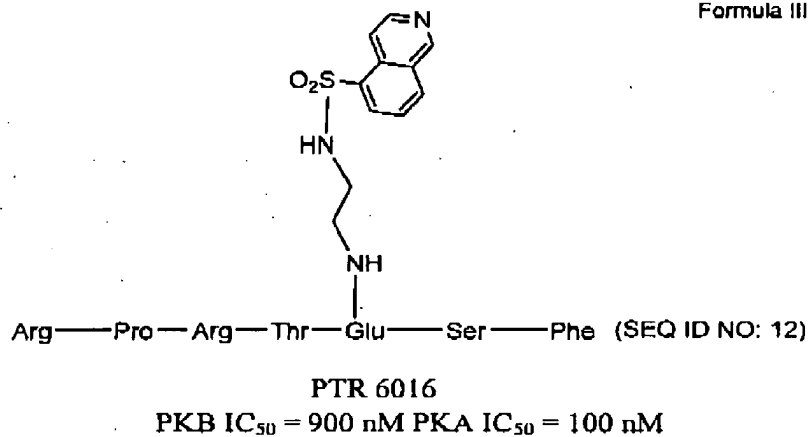
W is absent or is N-(8-sulfonamide-5-isoquinoline) ethylenediamine; and

L may be absent or is selected from the group consisting of glycine, β-alanine, phenylalanine, aminobutyric acid and aminopentanoic acid.

Preferably, W is connection through R<sub>5</sub> as described in formula IIb.

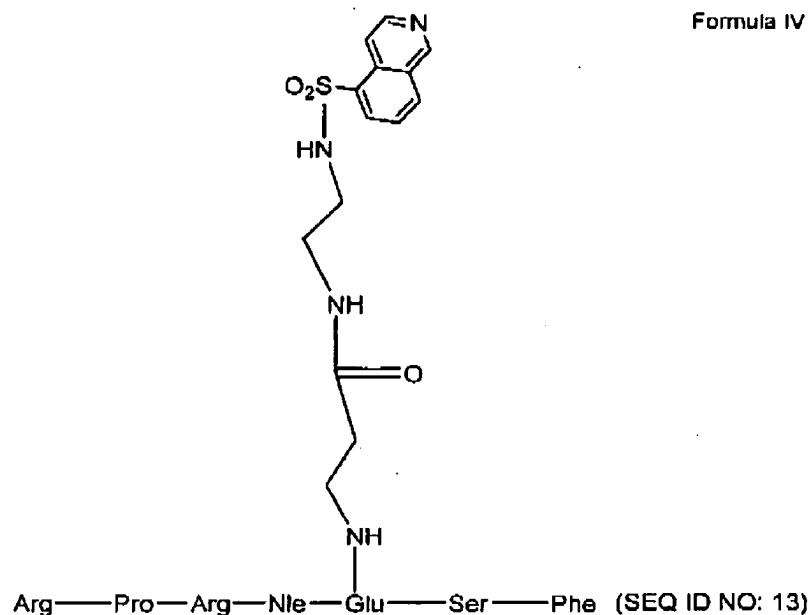
- 10 Currently most preferred embodiments of the present invention include a chimeric compound selected from the compounds described hereinbelow in formulae III-VII:

Formula III



15

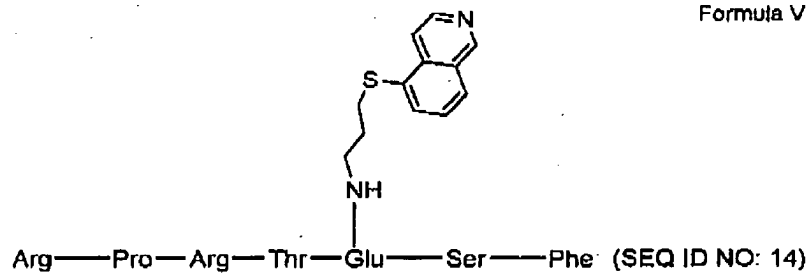
Formula IV



PTR 6086

PKB  $IC_{50}$  = 570 nM PKA  $IC_{50}$  = 500 nM

Formula V



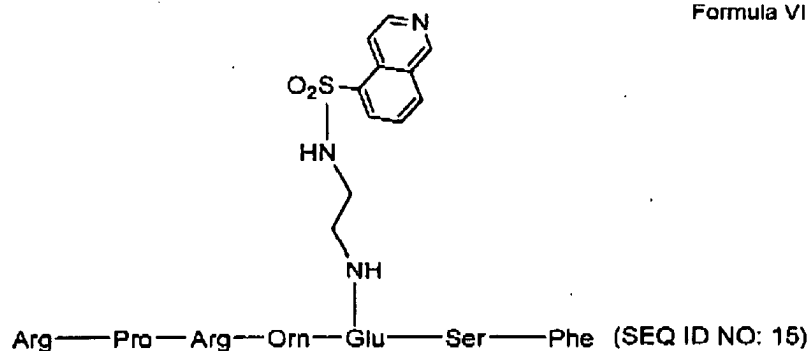
PTR 6104

PKB  $IC_{50}$  = 100 nM PKA  $IC_{50}$  = 100 nM

5



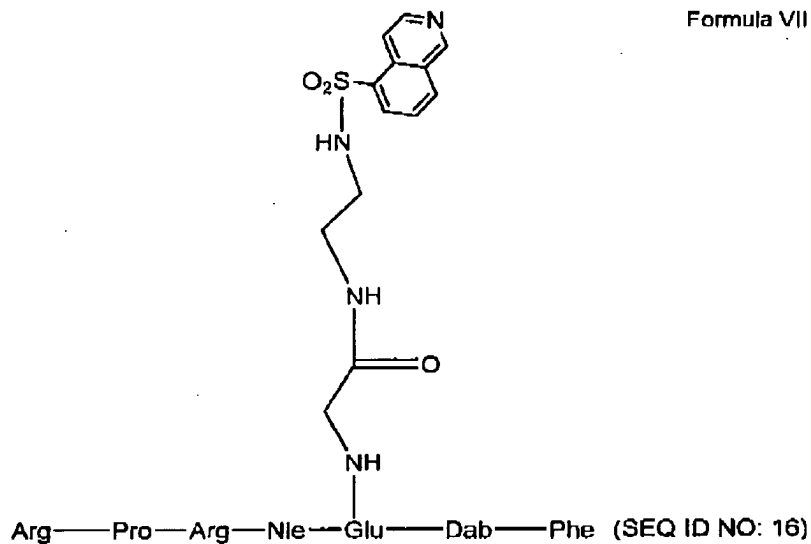
Formula VI



PTR 6132

PKB  $IC_{50}$  = 20 nM PKA  $IC_{50}$  = 12 nM

Formula VII



TY-60020-57

PKB  $IC_{50}$  = 70 nM PKA  $IC_{50}$  = 210 nM

The activity of the most preferred compounds according to formulae III-VII, as well as additional most preferred embodiments of the present invention, in inhibition of PKB and PKA activity is described in Table 1.

10 Currently most preferred are compounds comprising a sequence selected from:

Arg—Pro—Arg—Thr—Glu—(bAla-5-mercaptoaminopropyl-isoquinoline)—Ser—Phe  
(SEQ ID NO: 3).

Arg—Pro—Arg—Thr—Glu—(5-mercaptoaminopropyl-isoquinoline)—Ser—Phe (SEQ ID NO: 4).

Arg—Pro—Arg—Orn—Glu—(5-aminoethylsulfonamide-isoquinoline)—Ser—Phe (SEQ ID NO: 5).

5 Arg—Pro—Arg—Nva—Glu—(5-mercaptoaminopropyl-isoquinoline)—Ser—Phe (SEQ ID NO: 6).

Arg—Pro—Arg—Nle—Glu—(5-mercaptoaminopropyl-isoquinoline)—Ser—Phe (SEQ ID NO: 7).

10 Arg—Pro—Arg—Orn—Glu—(Gly-5-aminoethylsulfonamide)—Dab—Hol (SEQ ID NO: 8).

Arg—Pro—Arg—Nle—Glu—(Gly-5-aminoethylsulfonamide)—Dab—Phe (SEQ ID NO: 9).

Arg—Pro—Arg—Nle—Glu—(Gly-5-aminoethylsulfonamide)—Dab—Hol (SEQ ID NO: 10).

15

**Table 1. Most preferred compounds**

Compound	IC <sub>50</sub> PKB nM	IC <sub>50</sub> PKA nM
<b>PTR 6102</b>	300	100
<b>PTR 6104</b>	100	100
<b>PTR 6132</b>	20	12
<b>PTR 6134</b>	217	18
<b>PTR 6136</b>	114	11
<b>TY-60020-42</b>	110	50
<b>TY-60020-57</b>	70	210
<b>TY-60020-58</b>	120	50

20 Figure 1 describes the synergistic effect of conjugation of the ATP mimic part which poorly inhibit PKB, with the peptide substrate part having activity of 4  $\mu$ M, to yield the chimeric compound PTR 6016 with activity of 0.9  $\mu$ M activity.

Additional chimeric compounds comprising active peptides with diverse linkers connecting the peptidic moiety region to the spacer (according to scheme I). The favorable linker

enables simultaneous fit of both the peptide and ATP mimic into their substrate and ATP binding sites, and improvement of activity and specificity.

Additional preferred peptides according to the present invention may be used as the substrate domain of the chimeras and a basis for design of peptidomimetics. For example, linear  
5 7-mer peptides with  $IC_{50}$  of 4-5  $\mu$ M for PKB are disclosed in comparison to the reference 7-mer substrate that has  $K_m$  of 15  $\mu$ M. These peptides are specific to PKB and do not inhibit PKA activity at 60  $\mu$ M. Additional peptides disclosed demonstrate activity of 0.5-10  $\mu$ M for inhibition of PKB, and inhibit PKA at an  $IC_{50}$  of about 100 nM.

Peptidomimetic compounds having improved stability and cell permeability properties  
10 are another embodiment of the present invention. Non-limiting examples of generation of such compounds include N alkylation of selected peptide residues, carbamate, urea and hydrazine bond replacement, and incorporation of heterocyclic non-peptide moieties such as piperidine, piperazine, pyrrolidine etc. through peptide or non-peptide bond.

## 15 General methods

### Synthetic methods:

#### General methods for synthesis of peptides and carbamate bond formation.

The following procedure describes the synthesis of peptides in 96 wells plate (MPS plate) at a scale of 6  $\mu$ mol peptide per well, on Rink amide resin, using bis-(trichloromethyl)carbonate  
20 (BTC) for carbamate formation and HBTU/HOBT for normal coupling.

One gram of rink amide 0.6 mmol/g was swelled overnight in NMP with gentle shaking. The resin was distributed into 96 wells plate (~10 mg per well).

Fmoc deprotection performed by adding 500  $\mu$ l of 25% piperidine solution in NMP to each well and mixing at 650 rpm for 15 min, the piperidine solution is removed by a pressure of  
25 nitrogen and another portion of piperidine solution is added and shaken for 15 min. Wash of resin after Fmoc deprotection and after couplings, performed by placing 600  $\mu$ l NMP into each well, mixing for 2 min. and removing the NMP by nitrogen pressure. The washing procedure is repeated four times.

Regular coupling is performed by adding a solution of Fmoc protected amino acids (150  
30  $\mu$ l, 0.2M) in HOBT/NMP to the resin, followed by addition of HBTU solution in DMF (150  $\mu$ l,

0.2M) and DIEA in NMP (150  $\mu$ l, 0.4M). The reaction vessel block is mixed at 650 rpm for 1h and then removed by a pressure of nitrogen. This procedure is repeated once.

Carbamate formation using BTC is performed by addition of a solution of Fmoc protected amino alcohols in dioxane (150  $\mu$ l, 0.2 M), to a preactivation deep well plate, followed  
5 by addition of 150  $\mu$ l BTC (0.07 M) in 1,3-dichloropropane, 150  $\mu$ l collidine (0.6 M) in 1,3-dichloropropane and 200  $\mu$ l  $\text{CH}_2\text{Br}_2$ . The isocyanate solution is then transferred into the reaction vessel block and mixed for 40 min at 60°C. After 40 min the reaction mixture is removed by nitrogen pressure followed by  $\text{CH}_2\text{Cl}_2$  wash (3x 400  $\mu$ l). This procedure is repeated for additional two times.

10 Cleavage and global deprotection are performed by transferring the resin from the reaction vessel block into a deep well microtiter plate (cleavage plate). To this plate 350  $\mu$ l solution of 92.5% TFA, 2.5%  $\text{H}_2\text{O}$ , 2.5% TIS, 2.5% EDT is added. The plate is mixed at 1000 rpm for 1h and then the TFA solution is evaporated to dryness.

Purification by Sep-Pak performed by dissolving the residue of the resin with the peptide  
15 in 900  $\mu$ l solution A (0.1% TFA in water) and applying on C-18 Sep-Pak column. The peptides are eluted from the C-18 column by addition of 900  $\mu$ l solution A +  $\text{CH}_3\text{CN}$  1:1 to a deep well plate. The plate is frozen in liquid nitrogen at least 15 min and the peptides are lyophilized.

#### General methods for synthesis of chimeras in MPS format

20 The following procedure describes the synthesis of peptides in 96 wells plate (MPS plate) at a scale of 6  $\mu$ mol peptide per well, on Rink amide resin, using HBTU/HOBt for normal coupling.

One gram of rink amide 0.6 mmol/g was swelled overnight in NMP with gentle shaking. The resin was distributed into 96 wells plate (~10 mg per well).

25 Fmoc deprotection performed by adding 500  $\mu$ l of 25% piperidine solution in NMP to each well and mixing at 650 rpm for 15 min, the piperidine solution is removed by a pressure of nitrogen and another portion of piperidine solution is added and shaken for 15 min. Wash of resin after Fmoc deprotection and after couplings, performed by placing 600  $\mu$ l NMP into each well, mixing for 2 min. and removing the NMP by nitrogen pressure. The washing procedure is  
30 repeated four times.

Regular coupling is performed by adding a solution of Fmoc protected amino acids (150  $\mu$ l, 0.2 M) in HOBt/NMP to the resin, followed by addition of HBTU solution in DMF (150  $\mu$ l, 0.2 M) and DIEA in NMP (150  $\mu$ l, 0.4 M). The reaction vessel block is mixed at 650 rpm for 1h and then removed by a pressure of nitrogen. This procedure is repeated once. The last amino acid  
5 used in the assembly is N-Boc protected. At the end of assembly allyl deprotection takes place (from Glu(OAllyl) or C-building unit) by placing 500  $\mu$ l solution of Pd(Pphe3)<sub>4</sub> (0.02 M in chloroform containing 5% AcOH +2.5% NMM and mixing for 1h. This procedure is repeated once. Wash of the resin after allyl deprotection performed by addition of 600  $\mu$ l chloroform to each well and mixing for 5 min. The solvent is removed by nitrogen pressure. This wash is  
10 repeated for additional four times. The coupling of allyl protected linker to the peptide-resin is carried out by placing allyl protected linker (150  $\mu$ l, 0.2 M in NMP) followed by addition of PyBoP (0.2 M, in NMP) and DIEA (0.4 M, in NMP). The reaction vessel block is mixed for 1h the solution is removed by a pressure of nitrogen. This procedure is repeated once. The resin after the coupling is washed by addition of 500  $\mu$ l NMP to each well. Allyl removal from the  
15 linker is carried out followed the same procedure described above. After allyl deprotection, a solution of isoquinoline derivative (150  $\mu$ l, 0.2 M in NMP) is added followed by addition of ByBoP (150  $\mu$ l, 0.2 M in NMP) and DIEA (150  $\mu$ l, 0.4 M in NMP). The reaction block is mixed for 2 h.

Wash of the resin after this coupling performed by addition of 600 $\mu$ l NMP to each well  
20 and mixing for 2min. The solvent is removed by nitrogen pressure. This wash is repeated for additional four times.

Cleavage and global deprotection are performed by transferring the resin from the reaction vessel block into a deep well microtiter plate (cleavage plate). To this plate 350  $\mu$ l solution of 92.5% TFA, 2.5% H<sub>2</sub>O, 2.5% TIS, 2.5% EDT is added. The plate is mixed at 1000  
25 rpm for 1h and then the TFA solution is evaporated to dryness.

Purification by Sep-Pak performed by dissolving the residue of the resin with the peptide in 900  $\mu$ l solution A (0.1% TFA in water) + CH<sub>3</sub>CN 1:1 and applying on C-18 Sep-Pak column. This procedure is repeated once more. The plate is frozen in liquid nitrogen at least 15 min and the peptides are lyophilized.

30

## Biological screening assays for Inhibition of Protein Kinase activity:

### Cell free system methods

#### In vitro PKA kinase activity assay:

5 1. PKA enzyme was purchased from Promega. PKA activity is assayed on a 7-mer peptide, LRRASLG, known as kemptide. The assay is carried out in 96-well plates, in a final volume of 50  $\mu$ l per well. The reaction mixture includes various concentrations of the inhibitor, 50 mM MOPS, 10 mM MgAc, 0.2 mg/ml BSA, 10  $\mu$ M ATP, 20  $\mu$ M Kemptide and 1  $\mu$ Ci  $\gamma$ <sup>32</sup>P ATP. Reaction is started with addition of 15  $\mu$ l of the catalytic subunit of PKA diluted in 0.1  
10 mg/ml BSA, 0.4 U/well. Two blank wells without enzyme are included in every assay. The plates are agitated continuously at 30°C for 10' or at 27°C for 1 hour. Reaction is stopped by addition of 12  $\mu$ l 200 mM EDTA. 20  $\mu$ l aliquots of the assay mixture are spotted onto 2 cm<sup>2</sup> phosphocellulose strips (e.g. Whatman P81) and immersed in 75 mM phosphoric acid (10 ml per sample). The phosphocellulose strips are washed 6 times. Washes are done in continuous  
15 swirling for 5 minutes. last wash is in acetone. After air drying the strips, radiation is measured by scintillation spectrometry.

2. Screening compounds for PKA inhibition was performed in 96-well plate using SPA beads, as described below for PKB with the following modifications;  
The enzyme substrate was 5  $\mu$ M biotinylated -kemptide peptide (biotin-KLRRASLG). The  
20 kinase buffer was 50mM MOPS pH 7, 0.2 mg/ml BSA, 10mM Magnesium acetate. PKA (0.4 unit) diluted in 0.1 mg/ml BSA was added to each well.

#### **PKB in vitro kinase activity assays.**

1. PKB activity is assayed as described in Alessi et al. (FEBS Letters 399, 333, 1996)  
25 with the following modifications: instead of HA-PKB coupled to beads, soluble His-HA-PKB is used following precipitation on a Nickel column. The enzyme activity measurement is performed as described in the assay for PKA.

2. Screening compounds for PKB inhibition was performed in 96-well plate using method described previously (Kumar et al, BBA, 1526: 257-268, 2001) with modifications.  
30 Kinase reaction was carried out in final volume of 50  $\mu$ l. Each well contained 2.5  $\mu$ M of biotinylated-crosstide peptide (biotin-KGRPRTSFA) in kinase buffer [50 mM Tris-HCl pH 7.5, 10 mM MgCl<sub>2</sub>, 1 mM DTT and 0.1 mM sodium orthovanadate, 0.01% Triton X-100 and 2%

dimethyl ( $\text{Me}_2\text{SO}$ ), His-PKB enzyme and the potential inhibitory compound. The kinase reaction was started by adding  $10\mu\text{l}$  of  $2\mu\text{M}$  cold ATP and  $0.25\mu\text{Ci}$  of  $[\gamma^{33}\text{P}]\text{-ATP}$  in kinase buffer. The plates were incubated at  $27^\circ\text{C}$  for 1hr. At the end of the incubation the reaction was stopped by  $200\mu\text{l}$  of PBS containing 0.1% Triton X-100, 5 mM EDTA, 1 mM ATP and 0.3 mg/ml of Streptavidin-coated SPA bead (Amersham Pharmacia Biotech). After 15 min incubation at room temperature, the reaction mixtures were filtered using Packard GF/B 96-well plates. The plates were washed twice with 2M NaCl and 1% orthophosphoric acid followed by ethanol wash and 1h air-dry. The radioactivity was counted using microplate counter Packard Top Count.

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#### Transfer ELISA assay for measuring PKB activity and inhibition.

The inhibitor tested is dissolved in water to the desired concentration. Five  $\mu\text{l}$  of the inhibitor solution is added to the wells of a V shaped polypropylene microplate. Five  $\mu\text{l}$  of substrate peptide (Biotin—Lys—Gly—Arg—Pro—Arg—Thr—Ser—Ser—Phe—Ala—Glu—Gly (SEQ ID NO: 11)) solution in water at a concentration of  $300\mu\text{M}$  is then added to the wells (final assay concentration is  $100\mu\text{M}$ ). Then PKB enzyme dissolved in 3x reaction mixture (50 mM Tris HCl pH 7.5, 0.1% beta mercaptoethanol,  $1\mu\text{M}$  PKI (Calbiochem), 10 mM Mg acetate, ATP  $5\mu\text{M}$ ), is added in pre-calibrated amount to the wells. The amount of enzyme is calibrated so that less than 10% of the substrate is phosphorylated by the end of the reaction as evaluated by mass spectral analysis. The plate is covered with an adhesive tape, placed over a 1 mm ID vortex at  $30^\circ\text{C}$  and incubated for 30 min to 1 hour as needed. At the end of the incubation period  $5\mu\text{l}$  of 0.5 M disodium EDTA are added to the wells followed by  $180\mu\text{l}$  of PBS.

15

For ELISA, a microplate (Costar A/2) is coated with  $20\mu\text{l}$  of  $10\mu\text{g/ml}$  of avidin in PBS (over night at  $4^\circ\text{C}$  or 30 minutes at  $37^\circ\text{C}$ , on a 1 mm ID vortex). The plate is than washed several times with dionized water and flicked dry on a towel paper. The wells are filled with  $20\mu\text{l}$  of PBT (PBS + 1% BSA + 0.05% tween 20). Five  $\mu\text{l}$  from the enzyme reaction plate are transferred to the ELISA plate. The ELISA plate is placed on the 1 mm ID vortex and incubated for 10 min at RT. The plate is than washed with water as before. To each well  $20\mu\text{l}$  of anti phosphopeptide antibody (Cell Signaling Technology) diluted 1:1000 in PBT are added. The plate is placed again on the vortex, incubated for 30 minutes and washed with water as before. To each well  $20\mu\text{l}$  of goat anti- rabbit Ig conjugate with horse raddish peroxidase (HRP) is

25

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added. The plate is placed on the vortex, incubated for 20 min and washed with water as before. To each well is added 20  $\mu$ l of HRP substrate (Sigmafast OPD). After sufficient color development (up to maximum of about 30 minutes development time) the reaction is terminated by the addition of 20  $\mu$ l per well of 4 M HCl in water. The plate is then read using an ELISA reader at 490 nm. The signal obtained from wells containing potential inhibitors is compared to signal obtained from wells containing only the enzyme without inhibitor (maximum signal) and wells not containing enzyme (minimum signal).

The fraction of phosphorylated peptide can be also analyzed by mass spectra following desalting on a ziptip (C18, Millipore i). Mass of double charged substrate peptide is 759.3 Dalton, and of the double charged phosphorylated peptide is 799.3 Dalton.

#### PKC in vitro kinase assays

PKC was obtained from Promega Corp. and assayed according to the manufacturer's instructions using a kit from the same manufacturer, in the presence and absence of phospholipids. The activity of PKC was determined by subtracting the activity in the absence of phospholipids from that in the presence of phospholipids. The concentration of the ATP in the assay was 10 $\mu$ M ( $K_m$  for ATP=50 $\mu$ M).

#### Assays for inhibition of PKB activity in intact cells:

Several cancer cell lines were used to determine the activity of PKB inhibitors in intact cells. For example OVCAR3 is a cell line of ovarian carcinoma with an amplification of the PKB gene, U87MG is a glioma cell line with a deletion of PTEN gene - causing high activity of PKB, and PANC1 is a pancreatic carcinoma cell line with an amplification of PKB gene., PC-3, DU-145 and LNCaP are prostate cancer cell lines with alteration in PKB activity.

#### a. Annexin-V - apoptosis assay:

Cells were assayed for apoptosis using the Annexin-V (Bender medsystems). Cells were seeded in 6-well plates ( $0.3 \times 10^6$ /well) and treated with different concentrations of the inhibitors. At different time point, cells were scraped using rubber policemen, dispensed through syringe needle, washed twice with PBS and suspended in Annexin-V binding buffer (10mM Hepes/NaOH pH 7.4, 140mM NaCl and 2.5mM  $\text{CaCl}_2$ ). Annexin-V was diluted 1:40 and added



to each sample with 1 µg/ml Propidium Iodide (PI).  $0.5 \times 10^6$  cells were taken per sample for FACS analysis for apoptosis measurement.

- In an alternative method, cells were seeded in 10 cm plates ( $2 \times 10^6$  cells/plate) and treated with different concentrations of the inhibitor. 40 hours after treatment cells were trypsinised, washed twice with PBS and suspended in annexin-V buffer. annexin-V (Roche) is diluted 1:250 in a buffer containing 10 mM HEPES pH 7.4, 140 mM NaCl, 5 mM  $\text{CaCl}_2$  and 0.2 nM propidium iodid (PI). Apoptosis measurement was performed by FACS analysis.

b. ELISA assay for detection of ssDNA – apoptosis assay

- Cells were assayed for Apoptosis using the ssDNA Apoptosis ELISA kit (Chemicon International Inc). Cells were seeded in 96-well plates (5000 cells/well) and treated with different concentrations of the inhibitors. At different time point, plates were centrifuges at 200g for 5 min, medium was removed and the cells were fixed with 80% methanol in PBS for 30 min at room temperature. Plates were dried by floating in a waterbath at 37°C for 20 minutes. 50 µl of Formamide was added to each well and incubate at room temperature for 10 min. Plates were heated to 75°C in a circulating waterbath for 10 min, cooled in refrigerator for 5 min, and then formamide was removed. Plates were blocked by 3% non-fat dry milk in distilled water (w/v) for 1 hr at 37°C. Blocking was removed and 100 µl of antibody mixture (primary monoclonal antibody to ssDNA and HRP-labelled anti-mouse IgM) was added to each well. Plates were monitored for colour development by ABTS using microplate reader at 405 nm.

c. Cell viability assays:

- Cells were seeded in 96-well plates. After 72 hours in culture cells were treated with or without different concentrations (1, 5, 10, 25, 50, 100 µM) of the inhibitor, in triplicates for one to six days. Cells viability were tested using three methods: A. staining of viable cells with methylene blue, B. measurement the activity of mitochondrial dehydrogenases in viable cells using WST-1 reagent, c. incorporation of  $^3\text{H}$ -thymidine.

- Staining viable cells with methyelen blue: Cells were fixed by 0.5% gluterdialdehyde followed by staining with 1% methylene blue in borate buffer (Sigma) for one hour. Cells then washed few times with distilled water, air dried and the color was extracted by adding 0.1 M HCl for one hour at 37°C. Quantitation of color intensity was performed by measurement of the optical density at 620 nm by ELISA reader.

Cell proliferation reagent WST-1: At the appropriate time in culture medium was discarded and 100ul of WST-1 reagent (Boehringer mannheim) diluted 1:10 in growth medium was added for 1 – 2 hours at 37°C. The absorbency of the formazan product was measured at 450 nm with a reference wavelength of 690nm by microplate ELISA reader.

- 5 Incorporation of <sup>3</sup>H-thymidine: At the appropriate time in culture 1 uci of <sup>3</sup>H-thymidine (stock of 5 Ci/mmmole, Amersham) was added to each well containing 100ul of medium for 5 hours. At the end of the incubation the cells were washed few times with PBS, air dried for few hours and 50ul of scintillation liquid was added. The radioactivity was counted using microplate counter, Packard TopCount.

10

d. Inhibition of phosphorylation:

- Cells ( $2 \times 10^6$ ) were seeded in 25cm<sup>2</sup> flasks grown for 2 days at normal medium conditions then grown for additional 24 hours at starvation conditions (no FCS). At this time point, under starvation conditions, inhibitory compounds were added to analyze their effect on GSK-3 and PKB phosphorylation. At the end of the treatment cells were stimulated for 10 min with 150 ng/ml IGF-1 and lysed using lysis buffer (20 mM Tris-HCl pH 7.4, 150mM NaCl, 0.5% Triton-  
15 x100, 25mM NaF, 2mM AEBSF, 1mM sodium orthovanadate, 10mM  $\beta$ -glycerophosphate, 1 $\mu$ g/ml aprotinin and 5 $\mu$ g/ml leupeptin). Equal amounts of cell protein were resolved by 10% SDS-PAGE and electroblotted to PVDF membranes. Western blot analysis was performed using  
20 antibodies against phospho-Akt1 (Ser473), or (Thr 308) and phospho-GSK3 $\alpha$  (Ser21) were obtained from Cell Signaling Technology.

- In an alternative method, cells were seeded in 6-well plates, and treated with different concentrations of the inhibitor. Treatment was taken either under serum containing media or under starvation for different time periods. After treatment cells are stimulated for 10' with IGF-  
25 1 (HEK-293 and PANC1 cells) or EGF (OVCAR3 and U89MG cells). Cell lysates are prepared using boiled sample buffer. Western blot analysis with  $\alpha$  phospho-GSK3 showed decrease in GSK3 phosphorylation. The effect was also tested on GSK3 phosphorylation by expression of kinase-dead-PKB in HEK-293 cells.

### **In vivo models for evaluation the activity of PKB inhibitors**

The compounds of the present invention are tested for their affect on tumor growth and regression, in xenografts derived from cancerous cell lines such as:

1. Prostate cancer cells PC3, LNCAP and DU145 in nude mice;
- 5 2. Ovarian carcinoma cells (OVCAR) in nude mice;
3. Pancreatic cancer cells (PUNC1) in nude mice.

Briefly, the cells are implanted subcutaneously into the animals, and the tumors are allowed to grow to approximately 0.5 mm. The appropriate doses of compounds, which will be determined experimentally by acute toxicity studies, will be injected to the tumor at various stages of its growth. Injection at early stages will reflect the compound effect on tumor growth, 10 injection into an established tumor will determine its effect on regression. In addition, synergy studies are planned, where the compounds are injected into the tumor along with a known chemotherapy agent, to evaluate synergistic effects resulting from tumor increased sensitivity to chemotherapy due PKB inhibition leading to increased apoptosis.

15 The skilled artisan will appreciate that the following examples are merely illustrative and serve as non limitative exemplification of the principles of the present invention and that many variations and modifications are possible within the scope of the currently claimed invention as defined by the claims which follow.

20

### **EXAMPLES**

#### **Example 1. Screening PKA inhibitors for PKB inhibition**

Since there are no known inhibitors of PKB, the structural similarity between PKB and other protein kinases was used to screen commercially available inhibitors of other protein 25 kinases, e.g., PKA and PKC, for PKB inhibition. The preliminary screen was conducted in order to define some structural motifs in active compounds that would assist in the initial design of a combinatorial library of candidate compounds.

It should be noted, however, that though this approach is very useful for rapid identification of lead molecules, the molecules that are identified would possess inhibition 30 activity against other kinases as well. Thus, this approach dictates research directed not only at optimization of the inhibitory activity, but also, and perhaps most importantly,

specificity-oriented research. Namely, substantial efforts are actually directed at modifying the selectivity profile, in order to obtain a profile of selectivity or specificity towards PKB.

The screen yielded two compounds that inhibited PKB in the 2-3  $\mu$ M range. H-89, a known PKA inhibitor, was chosen to be the basic scaffold for the design of the first library,

5 based on its structure and on synthetic and specificity considerations.

H-89 was further optimized using rational design and parallel synthesis methods as described in Israeli application No. 136458. It was concluded that the 5-isoquinoline-sulfonamide-ethylenediamine core is essential for activity and replacement with any other core, either as a sulfonamide or a carboxamide derivative, eliminated activity. The substrate mimetic region C was also studied and it was concluded that this region could contain a hydrophobic or heterocyclic moiety or a peptide capable of binding PKB. Outline of these findings is given in the following example. These compounds served as basis for the design of the ATP mimetic moiety of the chimeric compounds according to the present invention.

15 **Example 2. Chimeric compounds having ATP and substrate mimetic sites**

Chimeric molecules are designed to combine an ATP- mimetic together with a substrate-mimetic regions connected via a bridge. These chimeric molecules can bind to both the catalytic site and to the substrate site of protein kinases at the same time and may have a synergistic effect that affords unique potency and selectivity properties. These compounds are designed to include an ATP-mimetic molecule, connected via various spacers to a substrate mimetic portion.

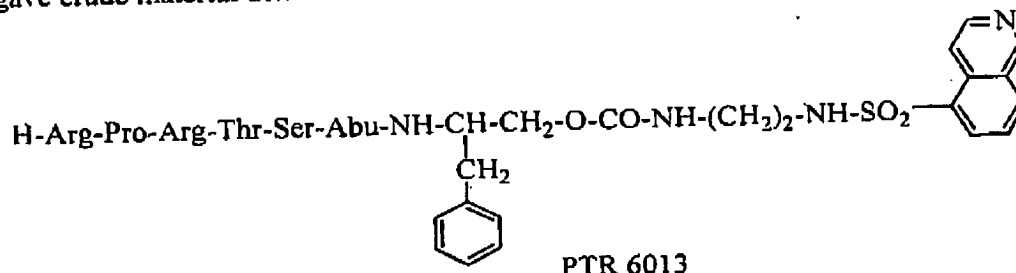
20 These compounds are identified following synthesis and screening cycles of combinatorial libraries in which each library examines modifications at a different region based on Scheme I, as shown above.

The ATP mimetic core includes but is not limited to dansyls, isoquinolines, quinolines and naphthalenes. The spacer is of varied lengths and conformations of any suitable chemistry including but not limited to amine, amide, thioether, oxyether, sulfonamide bond and the like. Non limiting examples for such spacers include sulfone amide derivatives, amino thiol derivatives and amino alcohol derivatives. The peptidic moiety comprises peptides and peptidomimetics. Such inhibitory peptides may be designed based on any peptide which may serve as a PKB substrate.

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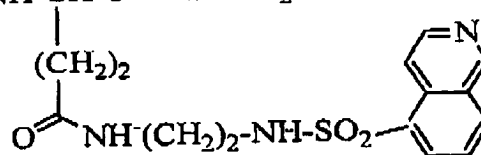
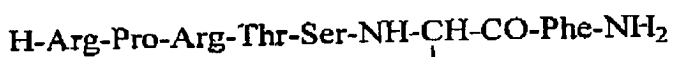
**Example 3. Detailed synthesis of chimeric compounds****PTR 6013**

Four hundred mg of 4-(4-formyl-3-methoxyphenoxy)butyryl (NovaGel HL) were swelled for 1.5 h in dichloroethane/trimethylorthoformate (1:1) in a reactor equipped with a sintered glass bottom, attached to a shaker. 352 mg (9 equivalents) of N-(8-sulfonamide-5-isoquinoline)ethylenediamine in 12 ml DMF was added to the resin followed by addition of NaBH(OAc)<sub>3</sub> and continues shaking over night. The resin was washed with DMF followed by DCM. Formation of the carbamate bond was performed by addition of Fmoc-Phenylalaninol (291 mg, 5 equivalents) which was preactivated with BTC (77 mg, 1.66 equivalents) and 2,4,6-collidine (290 µl, 14 equivalents) in THF twice at 50°C. Fmoc was removed from the resin using 25% Piperidine in NMP (3 ml) twice for 15 min followed by careful wash, seven times with NMP (5 ml), for 2 min each. Assembly of Abu, Ser, Thr, Arg, Pro Arg was accomplished by coupling cycles using Fmoc-Abu-OH, Fmoc-Ser(t-Bu)-OH, Fmoc-Thr(t-Bu)-OH, Fmoc-Arg(Pmc)-OH, and Fmoc-Pro-OH respectively. In each coupling cycle the amino acid (3 equivalents) was dissolved in NMP and was activated with PyBroP (3 equivalents) and DIEA (6 equivalents). Following coupling, the peptide-resin was washed, than Fmoc was removed followed by extensive wash with NMP, as described above for the first coupling. At the end of the assembly the peptide was cleaved from the resin using 65% TFA, 20% DCM, 5% thioanisole, 3% EDT, 2% TIS and 5% water in a total volume of 7 ml cocktail mixture for 15 min at 0°C under Argon and then 2h at room temperature. The solution was filtered through extract filter into polypropylene tube, the resin was washed with 3 ml of 60% TFA in DCM, the combined solution was evaporated by N<sub>2</sub> stream to give oily residue which on treatment with cold Et<sub>2</sub>O solidify. Centrifugation and decantation of the Et<sub>2</sub>O layer and treatment with additional portion of cold Et<sub>2</sub>O followed by centrifugation, decantation and drying of the white solid under vacuum over night, gave crude material denoted PTR 6013 having the following structure:



PTR 6014

Five hundred mg of Rink amide MBHA resin (0.55 mMol/g) were swelled for 2 h in NMP in a reactor equipped with a sintered glass bottom, attached to a shaker. Fmoc was removed from the resin using 25% Piperidine in NMP (4 ml) twice for 15 min followed by careful wash, seven times with NMP (5 ml), for 2 min each. Assembly of Phe, Glu, Ser, Thr, Arg, Pro, Arg was accomplished by coupling cycles using Fmoc-Phe-OH, Fmoc-Glu(Oallyl)-OH, Fmoc-Ser(t-Bu)-OH, Fmoc-Thr(t-Bu)-OH, Fmoc-Arg(Pmc)-OH, and Fmoc-Pro-OH respectively. In each coupling cycle, the amino acid (3 equivalents) was dissolved in NMP and was activated with PyBroP (3 equivalents) and DIEA (6 equivalents). At the end of assembly allyl deprotection took place using Pd(PPh<sub>3</sub>)<sub>4</sub> in solution of CH<sub>2</sub>Cl<sub>2</sub> containing 5% AcOH and 2.5% NMM. The free acid was activated by 3 equivalents PyBoP and 3.1 equivalents DIEA in NMP for 20 min followed by NMP wash. After preactivation a solution of small molecule (3 equivalents) and DIEA (4.5 equivalents) in NMP was added to the resin and shaken for 1h at room temperature. Following coupling, the peptide-resin was washed with NMP, than Fmoc was removed followed by extensive wash with NMP, as described above for the first coupling. At the end of the synthesis the peptide was cleaved from the resin using 85% TFA, 5% thioanisole, 3% EDT, 2% TIS and 5% water in a total volume of 5 ml cocktail mixture for 15 min at 0°C under Argon and then 2h at room temperature. The solution was filtered through extract filter into polypropylene tube, the resin was washed with 2 ml of TFA. The combined solution was evaporated by N<sub>2</sub> stream to give oily residue, which on treatment with cold Et<sub>2</sub>O solidifies. Centrifugation and decantation of the Et<sub>2</sub>O layer and treatment with additional portion of cold Et<sub>2</sub>O followed by centrifugation, decantation and drying the white solid under vacuum over night gave crude material denoted PTR 6014 having the following structure:

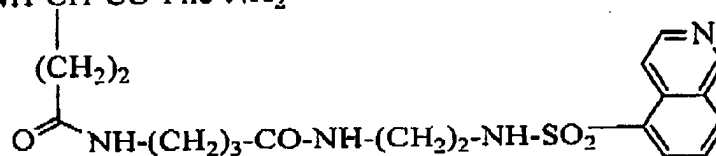
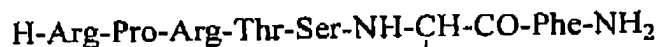


PTR 6014

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PTR 6020

Five hundred mg of Rink amide MBHA resin (0.55 mMol/g) were swelled for 2 h in NMP in a reactor equipped with a sintered glass bottom, attached to a shaker. Fmoc was removed from the resin using 25% piperidine in NMP (4 ml) twice for 15 min followed by careful wash, seven  
 5 times with NMP (5 ml), for 2 min each. Assembly of Phe, Glu, Ser, Thr, Arg, Pro, Arg was accomplished by coupling cycles using Fmoc-Phe-OH, Fmoc-Glu(Oallyl)-OH, Fmoc-Ser(t-Bu)-OH, Fmoc-Thr(t-Bu)-OH, Fmoc-Arg(Pmc)-OH, and Fmoc-Pro-OH respectively. In each coupling cycle the amino acid (3 equivalents) was dissolved in NMP and was activated with PyBroP (3 equivalents) and DIEA (6 equivalents). At the end of assembly allyl deprotection took place using  
 10  $\text{Pd(PPh}_3)_4$  in solution of  $\text{CH}_2\text{Cl}_2$  containing 5% AcOH and 2.5% NMM. The free acid was activated by 3 equivalents PyBOP and 3.1 equivalents DIEA in NMP for 20 min followed by NMP wash. After preactivation a solution of Allyl  $\gamma$ -aminobutyrate (5 equivalents) and DIEA (6 equivalents) in NMP was added and shaken for 1h at room temperature. The allyl deprotection and preactivation was performed by the same procedure as described above. A solution of the  
 15 small molecule (3 equivalents) and DIEA (4.5 equivalents) in NMP was added to the preactivated peptide-resin and shaken for 1h at room temperature. Following coupling, the peptide-resin was washed with NMP, than Fmoc was removed followed by extensive wash with NMP, as described above for the first coupling. At the end of the synthesis the peptide was cleaved from the resin using 85% TFA, 5% thioanisole, 3% EDT, 2% TIS and 5% water in a total of 5 ml cocktail  
 20 mixture for 15 min at  $0^\circ\text{C}$  under Argon and then 2h at room temperature. The solution was filtered through an extract filter into a polypropylene tube, the resin was washed with 2 ml of TFA, and the combined solution was evaporated by  $\text{N}_2$  stream to give an oily residue which on treatment with cold  $\text{Et}_2\text{O}$  solidifies. Centrifugation and decantation of the  $\text{Et}_2\text{O}$  layer and treatment with additional portion of cold  $\text{Et}_2\text{O}$  followed by centrifugation, decantation and drying the white solid  
 25 under vacuum over night gave crude material denoted PTR 6020 having the following structure:



PTR 6020

**Example 4. Biological activity of chimeric compounds**

Four chimeric compounds were screened for PKB inhibition activity. Table 2 describes their structure and inhibition activities. Similar to the compound denoted B-11-1 these compounds are not specific for PKA.

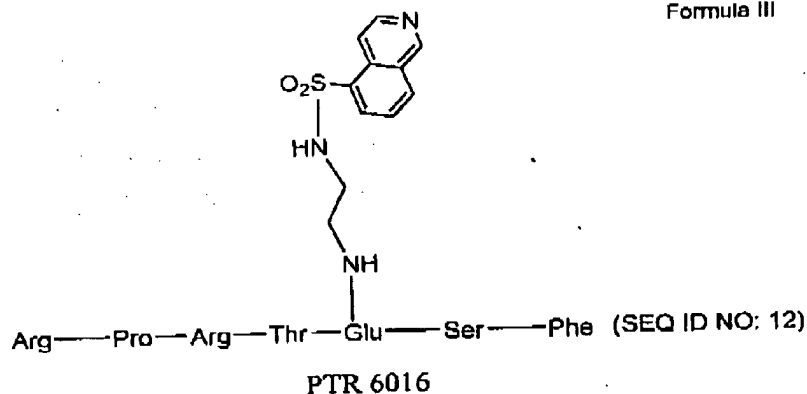
5

**Table 2.**

ID#	Activity PKB	Activity PKA
PTR 6013	3 $\mu$ M	1 $\mu$ M
PTR 6014	25 $\mu$ M	NA
PTR 6016	0.9 $\mu$ M	0.5 $\mu$ M
PTR 6020	> 20 $\mu$ M	NA

Structures of PTR 6013, 6014 and 6020 are described in example 11. The structure of PTR 6016 is:

Formula III



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Figure 1 describes the synergistic effect of conjugation of the ATP mimic part which does not inhibit PKB, with the peptide substrate part having activity of 4  $\mu$ M, to yield the chimeric compound PTR 6016 with activity of 0.9  $\mu$ M activity.

15

**Example 5. Additional chimeric compounds**

A multiple-parallel-synthesis of 96 chimeric compounds of active peptides from plates 60002, 60003 with diverse linkers (L in formulae II) connecting the peptide to the small molecule (W in Formulae II) was performed. These peptides were designed for elucidating the



appropriate linker which enables simultaneous fit of both the peptide and ATP mimic into their substrate and ATP binding sites, for improving activity and specificity.  
The compounds synthesized are described in formulae IIa-IId, as specified above.

## 5 Example 6. Peptides

Additional peptides are designed for use in the substrate domain of the chimeras and for design of peptidomimetics. Two plates of linear 7-mer peptides were synthesized and purified. Four peptides from the first plate (6002) were found to be active with  $IC_{50}$  of 4-5  $\mu$ M for PKB (the reference 7-mer substrate has  $IC_{50}$  of 15  $\mu$ M). All these peptides do not inhibit PKA activity at 60  $\mu$ M. From the second plate (6003) ~ 24 active peptides were identified, peptides exhibiting >60% inhibition were tested again, demonstrating activity of 0.5-10  $\mu$ M for PKB. Specificity to PKA not tested yet. Additional 1152 peptides from macrobeads library were screened, 150 had over 50% inhibition at 10  $\mu$ M. Three multiple-parallel-synthesis plates were then planed and synthesized. Selected results are presented in the following table.

15 Table 3.

	ID	structure	Activity	Activity
			PKB	PKA
20	TY 60002-50:	Arg-Pro-Arg-Thr-Ser-Ala-Hol (SEQ ID NO: 17)	5 $\mu$ M	>40 $\mu$ M
	TY 60002-61:	Arg-Pro-Arg-Val-Ser-Abu-Phe (SEQ ID NO: 18)	5 $\mu$ M	>40 $\mu$ M
	TY 60002-73:	Arg-Pro-Arg-Thr-Ser-Abu-Hol (SEQ ID NO: 19)	5 $\mu$ M	>40 $\mu$ M
	TY 60002-96:	Arg-Pro-Arg-Thr-Ser-Dap-Hol (SEQ ID NO: 20)	5 $\mu$ M	>40 $\mu$ M
	TY 60002-18:	Arg-Pro-Arg-Thr-Ser-Asp-Phe (SEQ ID NO: 21)	Not active	
25	AR 60003-50:	Arg-Pro-Arg-Met-Ser-Ser-Phe (SEQ ID NO: 22)	2.5 $\mu$ M	
	AR 60003-52:	Arg-Pro-Arg-Orn-Ser-Ser-Phe (SEQ ID NO: 23)	2.5 $\mu$ M	
	AR 60003-53:	Arg-Pro-Arg-Arg-Ser-Ser-Phe (SEQ ID NO: 24)	3 $\mu$ M	
	AR 60003-62:	Arg-Pro-Arg-Nle-Ser-Ser-Nle (SEQ ID NO: 25)	<1 $\mu$ M (70% inhibition at 1 $\mu$ M)	
	AR 60003-64:	Arg-Pro-Arg-Arg-ser-Ser-Arg (SEQ ID NO: 26)	Not active	
30	AR 60003-96:	Arg-Pro-Arg-Orn-Ala-Thr-Orn (SEQ ID NO: 27)	Not active	

The PKB inhibition activity of peptide AR-60003-52 as determined in ELISA is  
35 illustrated in figure 2.

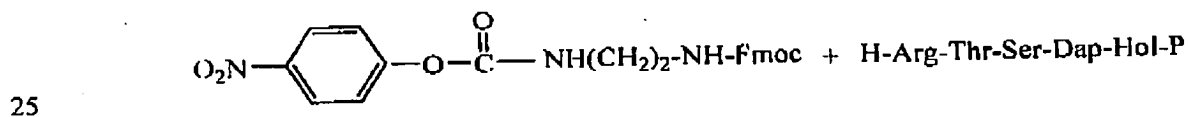
**Example 7. Peptidomimetic compounds based on the active peptides:**

The following peptidomimetic compounds which contain carbamate and/or urea bonds replacing peptidic bonds, were synthesized.

- 5 PTR 6046: H-Arg-NH-(CH<sub>2</sub>)<sub>2</sub>-NH-CO-Arg-Thr-Ser-Dap-Hol-NH<sub>2</sub>  
 PTR 6048: H-Arg-NH-CH<sub>2</sub>-C<sub>6</sub>H<sub>4</sub>-CH<sub>2</sub>-NH-CO-Arg-Thr-Ser-Dap-Hol-NH<sub>2</sub>  
 PTR 6050: H<sub>2</sub>N-(CH<sub>2</sub>)<sub>4</sub>-NH-CO-Pro-Arg-Thr-Ser-Dap-Hol-NH<sub>2</sub>  
 PTR 6052: H<sub>2</sub>N-(CH<sub>2</sub>)<sub>4</sub>-NH-CO-3-(HNCH<sub>2</sub>)-C<sub>6</sub>H<sub>4</sub>-CH<sub>2</sub>NH-CO-Arg-Thr-Ser-Dap-Hol-NH<sub>2</sub>  
 PTR 6054: H-Arg-OI-NH-CH<sub>2</sub>-C<sub>6</sub>H<sub>4</sub>-CH<sub>2</sub>-NH-CO-Arg-Thr-Ser-Dap-Hol-NH<sub>2</sub> (this compound  
 10 contains both carbamate and urea bonds).  
 PTR 6056: H-Arg-1,4-Homopiperazine-CO-Arg-Thr-Ser-Dap-Hol-NH<sub>2</sub>

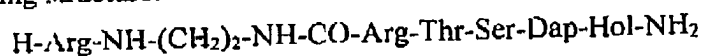
**Example 8. Detailed synthesis of PTR 6046, an heptamer containing an urea bond**

- One hundred mg (0.055 μmol) of rink amide resin were swelled 1.5h in NMP in a reactor  
 15 with a sintered glass bottom, attached to a shaker. Fmoc was removed from the resin using 25%  
 piperidine in NMP (3 ml) twice for 15 min followed by careful wash, seven times with NMP (2  
 ml). Assembly of Arg-Thr-Ser-Dap-Hol was accomplished by coupling cycles using Fmoc-Hol-  
 OH, Fmoc-Dap-OH, Fmoc-Ser(tBu)-OH, Fmoc-Thr(tBu)-OH, Fmoc-Arg(Pmc)-OH. In each  
 coupling cycle the amino acid (3 equivalents) was dissolved in NMP and was activated with  
 20 PyBroP (3 equivalents) and DIEA (6 equivalents). Following coupling, the peptide-resin was  
 washed, than Fmoc was removed followed by extensive wash with NMP.

Formation of the urea bond:

- 50 mg (2 equivalents) of N-Fluorenylmethoxycarbonyl-N'-nitrophenoxycarbonyl-  
 diaminoethane, 25 μl (2.5 equivalents) of DIEA in 2 ml NMP was added to the resin and  
 continued shaking 1.5h. The resin was washed with NMP (5 times 2 min each). After formation  
 30 of the urea bond, a coupling of Fmoc-Arg(Pmc)-OH was performed as describe above followed  
 by Fmoc deprotection. At the end of the assembly the peptide was cleaved from the resin using

92.5% TFA, 2.5% EDT, 2.5% TIS and 2.5% water in a total volume of 5 ml cocktail mixture and continues shaking 1 h. The solution was filtered through extract filter into polypropylene tube, the resin was washed with 2 ml of TFA, the combined solution was evaporated by N<sub>2</sub> stream to give oily residue which on treatment with cold Et<sub>2</sub>O solidify. Centrifugation and decantation of the Et<sub>2</sub>O layer and treatment with additional portion of cold Et<sub>2</sub>O followed by centrifugation and decantation and drying the white solid under vacuum over night gave crude PTR 6046 having the following structure:



10 **Example 9. Synthesis and screening of 96 chimeric compounds in MPS format**

The following sequences (denoted TY-60020- and shown in Table 4) were synthesized in MPS format according to the synthesis method above.

**Table 4.** Structure of compounds in TY-60020 plate

	1	2	3	4	5	Linker*	ATP mimetic	6	7
1	Arg	Pro	Arg	Dap	Glu	0	5-aminoethylsulfonamide isoquinoline	Dab	Phe
2	Arg	Pro	Arg	Dap	Glu	0	5-aminoethylsulfonamide isoquinoline	Dab	Hol
3	Arg	Pro	Arg	Dap	Glu	0	5-aminoethylsulfonamide isoquinoline	Orn	Phe
4	Arg	Pro	Arg	Dap	Glu	0	5-aminoethylsulfonamide isoquinoline	Orn	Hol
5	Arg	Pro	Arg	Dap	Glu	0	5-aminoethylsulfonamide isoquinoline	Ala	Phe
6	Arg	Pro	Arg	Dap	Glu	0	5-aminoethylsulfonamide isoquinoline	Ala	Hol
7	Arg	Pro	Arg	Dap	Glu	0	5-aminoethylsulfonamide isoquinoline	Nva	Phe
8	Arg	Pro	Arg	Dap	Glu	0	5-aminoethylsulfonamide isoquinoline	Nva	Hol
9	Arg	Pro	Arg	Orn	Glu	0	5-aminoethylsulfonamide isoquinoline	Dab	Phe
10	Arg	Pro	Arg	Orn	Glu	0	5-aminoethylsulfonamide isoquinoline	Dab	Hol
11	Arg	Pro	Arg	Orn	Glu	0	5-aminoethylsulfonamide isoquinoline	Orn	Phe
12	Arg	Pro	Arg	Orn	Glu	0	5-aminoethylsulfonamide isoquinoline	Orn	Hol

13	Arg	Pro	Arg	Orn	Glu	0	5-aminoethylsulfonamide isoquinoline	Ala	Phe
14	Arg	Pro	Arg	Orn	Glu	0	5-aminoethylsulfonamide isoquinoline	Ala	Hol
15	Arg	Pro	Arg	Orn	Glu	0	5-aminoethylsulfonamide isoquinoline	Nva	Phe
16	Arg	Pro	Arg	Orn	Glu	0	5-aminoethylsulfonamide isoquinoline	Nva	Hol
17	Arg	Pro	Arg	Abu	Glu	0	5-aminoethylsulfonamide isoquinoline	Dab	Phe
18	Arg	Pro	Arg	Abu	Glu	0	5-aminoethylsulfonamide isoquinoline	Dab	Hol
19	Arg	Pro	Arg	Abu	Glu	0	5-aminoethylsulfonamide isoquinoline	Orn	Phe
20	Arg	Pro	Arg	Abu	Glu	0	5-aminoethylsulfonamide isoquinoline	Orn	Hol
21	Arg	Pro	Arg	Abu	Glu	0	5-aminoethylsulfonamide isoquinoline	Ala	Phe
22	Arg	Pro	Arg	Abu	Glu	0	5-aminoethylsulfonamide isoquinoline	Ala	Hol
23	Arg	Pro	Arg	Abu	Glu	0	5-aminoethylsulfonamide isoquinoline	Nva	Phe
24	Arg	Pro	Arg	Abu	Glu	0	5-aminoethylsulfonamide isoquinoline	Nva	Hol
25	Arg	Pro	Arg	Nle	Glu	0	5-aminoethylsulfonamide isoquinoline	Dab	Phe
26	Arg	Pro	Arg	Nle	Glu	0	5-aminoethylsulfonamide isoquinoline	Dab	Hol
27	Arg	Pro	Arg	Nle	Glu	0	5-aminoethylsulfonamide isoquinoline	Orn	Phe
28	Arg	Pro	Arg	Nle	Glu	0	5-aminoethylsulfonamide isoquinoline	Orn	Hol
29	Arg	Pro	Arg	Nle	Glu	0	5-aminoethylsulfonamide isoquinoline	Ala	Phe
30	Arg	Pro	Arg	Nle	Glu	0	5-aminoethylsulfonamide isoquinoline	Ala	Hol
31	Arg	Pro	Arg	Nle	Glu	0	5-aminoethylsulfonamide isoquinoline	Nva	Phe
32	Arg	Pro	Arg	Nle	Glu	0	5-aminoethylsulfonamide isoquinoline	Nva	Hol
33	Arg	Pro	Arg	Dap	Glu	Gly	5-aminoethylsulfonamide isoquinoline	Dab	Phe
34	Arg	Pro	Arg	Dap	Glu	Gly	5-aminoethylsulfonamide isoquinoline	Dab	Hol

35	Arg	Pro	Arg	Dap	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Orn	Phe
36	Arg	Pro	Arg	Dap	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Orn	Hol
37	Arg	Pro	Arg	Dap	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Ala	Phe
38	Arg	Pro	Arg	Dap	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Ala	Hol
39	Arg	Pro	Arg	Dap	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Nva	Phe
40	Arg	Pro	Arg	Dap	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Nva	Hol
41	Arg	Pro	Arg	Orn	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Dab	Phe
42	Arg	Pro	Arg	Orn	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Dab	Hol
43	Arg	Pro	Arg	Orn	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Orn	Phe
44	Arg	Pro	Arg	Orn	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Orn	Hol
45	Arg	Pro	Arg	Orn	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Ala	Phe
46	Arg	Pro	Arg	Orn	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Ala	Hol
47	Arg	Pro	Arg	Orn	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Nva	Phe
48	Arg	Pro	Arg	Orn	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Nva	Hol
49	Arg	Pro	Arg	Abu	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Dab	Phe
50	Arg	Pro	Arg	Abu	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Dab	Hol
51	Arg	Pro	Arg	Abu	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Orn	Phe
52	Arg	Pro	Arg	Abu	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Orn	Hol
53	Arg	Pro	Arg	Abu	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Ala	Phe
54	Arg	Pro	Arg	Abu	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Ala	Hol
55	Arg	Pro	Arg	Abu	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Nva	Phe
56	Arg	Pro	Arg	Abu	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Nva	Hol

57	Arg	Pro	Arg	Nle	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Dab	Phe
58	Arg	Pro	Arg	Nle	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Dab	Hol
59	Arg	Pro	Arg	Nle	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Orn	Phe
60	Arg	Pro	Arg	Nle	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Orn	Hol
61	Arg	Pro	Arg	Nle	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Ala	Phe
62	Arg	Pro	Arg	Nle	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Ala	Hol
63	Arg	Pro	Arg	Nle	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Nva	Phe
64	Arg	Pro	Arg	Nle	Glu	Gly	5-aminoethylsulfoneamide isoquinoline	Nva	Hol
65	Arg	Pro	Arg	Dap	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Dab	Phe
66	Arg	Pro	Arg	Dap	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Dab	Hol
67	Arg	Pro	Arg	Dap	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Orn	Phe
68	Arg	Pro	Arg	Dap	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Orn	Hol
69	Arg	Pro	Arg	Dap	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Ala	Phe
70	Arg	Pro	Arg	Dap	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Ala	Hol
71	Arg	Pro	Arg	Dap	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Nva	Phe
72	Arg	Pro	Arg	Dap	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Nva	Hol
73	Arg	Pro	Arg	Orn	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Dab	Phe
74	Arg	Pro	Arg	Orn	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Dab	Hol
75	Arg	Pro	Arg	Orn	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Orn	Phe
76	Arg	Pro	Arg	Orn	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Orn	Hol
77	Arg	Pro	Arg	Orn	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Ala	Phe
78	Arg	Pro	Arg	Orn	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Ala	Hol

79	Arg	Pro	Arg	Orn	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Nva	Phe
80	Arg	Pro	Arg	Orn	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Nva	Hol
81	Arg	Pro	Arg	Abu	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Dab	Phe
82	Arg	Pro	Arg	Abu	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Dab	Hol
83	Arg	Pro	Arg	Abu	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Orn	Phe
84	Arg	Pro	Arg	Abu	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Orn	Hol
85	Arg	Pro	Arg	Abu	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Ala	Phe
86	Arg	Pro	Arg	Abu	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Ala	Hol
87	Arg	Pro	Arg	Abu	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Nva	Phe
88	Arg	Pro	Arg	Abu	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Nva	Hol
89	Arg	Pro	Arg	Nle	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Dab	Phe
90	Arg	Pro	Arg	Nle	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Dab	Hol
91	Arg	Pro	Arg	Nle	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Orn	Phe
92	Arg	Pro	Arg	Nle	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Orn	Hol
93	Arg	Pro	Arg	Nle	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Ala	Phe
94	Arg	Pro	Arg	Nle	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Ala	Hol
95	Arg	Pro	Arg	Nle	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Nva	Phe
96	Arg	Pro	Arg	Nle	Glu	bAla	5-aminoethylsulfoneamide isoquinoline	Nva	Hol

\* The ATP mimetic is connected via the specified linker to the carboxy group of the glutamic acid side chain at position 5.

The 96 compounds were screened at 1  $\mu$ M compound concentration (according to the above methods) for PKB and PKA activity inhibition (% inhibition of kinase activity) and the results are summarized in Table 5:

Table 5. Screening results of TY-60020 compounds

Sample #	PKB-kinase (spa)		PKA-kinase	
	Assay 1	Assay 2	Assay 1	Assay 2
60020-1	67	65	98	99
60020-2	47	48	94	95
60020-3	52	60	96	96
60020-4	34	53		
60020-5	46	26		
60020-6	32	14		
60020-8	18	0		
60020-9	71	57	96	97
60020-10	45	48	92	94
60020-11	58	52	95	96
60020-12	23	3		
60020-13	36	29		
60020-14	26	31		
60020-15	31	44		
60020-17	34	38		
60020-18	18	28		
60020-19	32	26		
60020-20	11	15		
60020-21	-2	6		
60020-22	-5	-2		
60020-23	-3	-8		
60020-24	-17	-12		
60020-26	17	18		
60020-27	63	25		
60020-28	27	31		
60020-29	19	39		
60020-30	14	38		
60020-32	12	32		
60020-33	63	69	98	98
60020-34	64	65	97	98
60020-35	43	56	95	96
60020-36	69	71	93	94
60020-37	53	44	98	99
60020-38	49	49	98	99
60020-39	17	16		
60020-40	36	22		
60020-42	80	86	99	99
60020-43	48	62	96	96
60020-44	54	67	95	95
60020-45	29	53		
60020-46	37	57		



60020-47	21	33		
60020-48	10	22		
60020-49	66	66	95	95
60020-50	60	54	97	97
60020-51	47	37	94	95
60020-52	56	53	92	92
60020-53	20	13		
60020-54	34	45		
60020-55	20	40		
60020-56	20	47		
60020-57	92	93	95	96
60020-58	89	93	98	99
60020-59	62	75	92	93
60020-60	59	70	87	89
60020-61	29	40		
60020-62	29	38		
60020-63	17	19		
60020-64	25	26		
60020-65	32	30		
60020-66	52	63		
60020-67	51	68		
60020-68	61	74		
60020-69	36	57		
60020-70	57	68		
60020-71	32	46		
60020-72	31	52		
60020-73	16	35		
60020-74	33	41		
60020-75	23	35		
60020-76	38	38		
60020-77	13	15		
60020-78	21	24		
60020-80	13	35		
60020-16	5	37		
60020-81	-17	42		
60020-82	-9	56		
60020-83	-21	60		
60020-84	10	51		
60020-85	-27	-2		
60020-86	-55	-5		
60020-87	-50	5		
60020-88	-50	31		
60020-89	8	48		
60020-90	19	27		
60020-92	40	59		
60020-93	-12	58		

60020-94	9	31		
60020-95	46	90		
60020-96	0	54		

Table 6. most active compounds from plate TY-60020

Compound	IC <sub>50</sub> PKB nM	IC <sub>50</sub> PKA nM
TY-60020-57	70	210
TY-60020-58	120	50
TY-60020-42	110	50
TY-60020-95	3300	2000
TY-60020-67	1080	250
TY-60020-68	500	250
TY-60020-59	300	200

**Example 10. Synthesis of 19 chimeric compounds**

- 5 Additional 19 compound were synthesized (denoted BP-60023- ) comprising N<sup>α</sup>-ω-functionalized derivative of an amino acid at position 5 incorporated in order to adjust the spatial position of the peptide and the small molecule relative to each other, by attachment to N-derivatized amino acid rather than to amino acid side chain. The sequences are shown in Table 7.

Table 7. MPS plate BP-60023

		1	2	3	4	5	6	7
1	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Nva	DTyrC4*	Dap	Hol
2	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	DTyrC4*	Dap	Hol
3	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Nva	TyrC4*	Dap	Hol
4	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	LTyrC4*	Dap	Hol
5	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	GlyC2*	Ser	Phe
6	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	GlyC3*	Ser	Phe
7	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	GlyC5*	Ser	Phe
8	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Nle	GlyC2*	Dab	Phe
9	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Nle	GlyC3*	Dab	Phe
10	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Nle	GlyC5*	Dab	Phe
11	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	GlyC2*	Dab	Phe
12	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	GlyC3*	Dab	Phe
13	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	GlyC5*	Dab	Phe
14	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	GlyC2*	Dab	Hol
15	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	GlyC3*	Dab	Hol
16	CO-N-SO <sub>2</sub> -IQ	Arg	Pro	Arg	Orn	GlyC5*	Dab	Hol

17	CO-N-SO2-IQ	Arg	Pro	Arg	Nva	GlyC2*	Ala	Hol
18	CO-N-SO2-IQ	Arg	Pro	Arg	Nva	GlyC3*	Ala	Hol
19	CO-N-SO2-IQ	Arg	Pro	Arg	Nva	GlyC5*	Ala	Hol

SO2-IQ = 5-aminoethylsulfoneamide isoquinoline

### Example 11. Structure and activity of chimeric PTRs

5

Table 8.

PTR	Structure	PKB IC <sub>50</sub> (uM)	PKA IC <sub>50</sub> (uM)
6013	Arg-Pro-Arg-Thr-Ser-Abu-Phe-(O-CO-N-SO2-IQ) (SEQ ID NO: 28)	4	na
6014	Arg-Pro-Arg-Thr-Ser-Glu-(CO-N-SO2-IQ)-Phe (SEQ ID NO: 29)	25	na
6016	Arg-Pro-Arg-Thr-Glu-(CO-N-SO2-IQ)-Ser-Phe (SEQ ID NO: 12)	1	0.1
6020	Arg-Pro-Arg-Thr-Ser-Abu-Phe-O-CO-N-bAla-SO2-IQ) (SEQ ID NO: 30)	18	na
6082	Arg-Pro-Arg-Thr-Glu-(CO-N-bAla-SO2-IQ)-Ser-Phe (SEQ ID NO: 31)	5.54	0.2
6086	Arg-Pro-Arg-Nle-Glu-(CO-N-bAla-SO2-IQ)-Ser-Phe (SEQ ID NO: 13)	0.57	0.5
6088	Arg-Pro-Arg-Orn-Glu-(CO-N-Gaba-SO2-IQ)-Ser-Phe (SEQ ID NO: 32)	1.18	1
6090	Arg-Pro-Arg-Thr-Glu-(CO-N-Ape5-SO2-IQ)-Ser-Phe (SEQ ID NO: 33)	5.6	0.6
6096	Arg-Pro-Arg-Nle-Glu-(CO-N-bAla-SO2-IQ)-Ser-Nle (SEQ ID NO: 34)	0.93	0.2
6102	Arg-Pro-Arg-Thr-Glu-(CO-N-bAla-S-IQ)-Ser-Phe (SEQ ID NO: 35)	0.3	0.1
6104	Arg-Pro-Arg-Thr-Glu-(CO-N-S-IQ)-Ser-Phe (SEQ ID NO: 14)	0.1	0.1
6106	Arg-Pro-Arg-Thr-Dap-(N-CO-SO2-IQ)-Ser-Phe # (SEQ ID NO: 36)	10	na
6128	Arg-Pro-Arg-Nle-Glu-(CO-N-SO2-IQ)-Ser-Phe (SEQ ID NO: 37)	9	0.1
6130	Arg-Pro-Arg-Thr-Asp-(CO-N-SO2-IQ)-Ser-Phe (SEQ ID NO: 38)	9.5	0.053
6132	Arg-Pro-Arg-Orn-Glu-(CO-N-SO2-IQ)-Ser-Phe (SEQ ID NO: 15)	0.02	0.012
6134	Arg-Pro-Arg-Nva-Glu-(CO-N-S-IQ)-Ser-Phe (SEQ ID NO: 39)	0.217	0.018
6136	Arg-Pro-Arg-Nle-Glu-(CO-N-S-IQ)-Ser-Phe (SEQ ID NO: 40)	0.114	0.011

6138	Arg-Pro-Arg-Nle-Glu-(CO-N-bAla-SO <sub>2</sub> -IQ)-Abu-Phe (SEQ ID NO: 41)	2	na
6140	Arg-Pro-Arg-Nle-Glu-(CO-N-bAla-SO <sub>2</sub> -IQ)-Dab-Phe (SEQ ID NO: 42)	0.413	0.195
6150	Arg-Pro-Arg-Nva-Glu-(CO-N-SO <sub>2</sub> -IQ)-Ala-Hol (SEQ ID NO: 43)	25	na
6152	Arg-Pro-Arg-Nva-Glu-(CO-N-bAla-SO <sub>2</sub> -IQ)-Ala-Hol (SEQ ID NO: 44)	5	5

# the small molecule of this compound is connected to the peptide via an amide bond between an amine of Dap residue on the peptide and the carboxylic acid on the small molecule, while in the other compounds the small molecule is connected to the peptide via an amide bond between a carboxylic moiety on the peptide and amino moiety on the small molecule.

5 na = not assayed

SO<sub>2</sub>-IQ = 5-aminoethylsulfonamide isoquinoline

S-IQ = 5-mercaptoaminopropyl isoquinoline

#### 10 Example 12. Selected peptides and peptidomimetic compounds serving as substrate-mimetic inhibitors

Several peptide and peptidomimetic compounds plates were screened for PKB and PKA inhibition. (These results relate to the peptide moiety alone, without the conjugation to a small molecule). Selected results are presented in Table 9.

15 Table 9.

Compound	Sequence							% Inhibition		
								PKB I	PKB II	PKA
60005-5	Arg	Pro	Arg	Thr	Ser	GlyNH <sub>2</sub>	Phe	96	93	40
60005-32	Arg	Pro	Arg	Thr	GlyNH <sub>2</sub>	Ala	Hol	43	44	4
60005-75	Arg	Pro	Arg	Nval	Ser	GlyNH <sub>2</sub>	Hol	70	70	-4
60005-84	Arg	Pro	Arg	Nval	Thr	Ala	Hol	39	48	7
60006-5A	Arg	Pro	Arg	Nval	GlyNH <sub>2</sub>	Ala	Hol	47	43	10
60006-7B	Arg	Pro	Arg	Nval	GlyNH <sub>2</sub>	Abu	Hol	53	47	24
60006-11B	Arg	Pro	Arg	Nval	GlyNH <sub>2</sub>	Dap	Ph <sub>2</sub>	51	44	20
60006-16A	Arg	Pro	Arg	Nval	GlyNH <sub>2</sub>	GltNH <sub>2</sub>	Hol	42	42	21
60006-18A	Arg	Pro	Arg	Nval	Gly	Ala	Hol	47	52	11
60002-38	Arg	Pro	Arg	Nval	Ser	Ala	Phe	43		-2
60002-50	Arg	Pro	Arg	Thr	Ser	Ala	Hol	62		2
60002-73	Arg	Pro	Arg	Thr	Ser	Abu	Hol	44		8
60002-83	Arg	Pro	Arg	Thr	Abu	Dap	Phe	40		-3

The most promising peptides from the above described screening were assayed for an accurate  $IC_{50}$  curve. In addition, several PTR compounds were designed and based on the screening results, and assayed as well. The results of the  $IC_{50}$  evaluation are compiled in Table 10.

Table 10.

	Compound								
IC50 μM	60002- 73	60002- 38	60006- 18A	60006- 8A	60006- 5A	60005- 5	60018- 16	PTR 6154	PTR 6158
PKA	228	80	>100	39	139	7.5	2	1.78	0.99
PKB	34	18	9.45	23	10	2.4	100	>50	>100

Figure 3 describes PKB and PKA inhibition curves ( $IC_{50}$ ) of the active and selective peptide denoted 60018-16.

Based on the best peptide PTR 6158 several analogs were synthesized bearing hydrophobic moieties, as depicted in Table 11. In addition, 5 compounds denoted BP60023 20-24 were synthesized comprising the peptide sequences in Table 11. These are peptides with hydrophobic moieties, aimed for improvement of cell permeability.

Peptides alone showed activity in the order of magnitude of 1  $\mu M$ , with selectivity PKB/PKA 100X. The peptide alone binds to the substrate site. When conjugated to an ATP moiety, the ATP binds first and the peptide is dislocated, thus the affinity is higher and the selectivity is lower. Synergistic effect on activity. ATP moiety does not distinguish between PKA/PKB sites, thus the ATP domain is not selective

Table 11.

20	Lauryl	Gly	Arg	Pro	Arg	Nva	Tyr	Dap	Hol
21	Lauryl	bAla	Arg	Pro	Arg	Nva	Tyr	Dap	Hol
22		Myristyl	Arg	Pro	Arg	Nva	Tyr	Dap	Hol
23	Myristyl	Gly	Arg	Pro	Arg	Nva	Tyr	Dap	Hol
24	Myristyl	bAla	Arg	Pro	Arg	Nva	Tyr	Dap	Hol

### Example 13. Induction of apoptosis in prostate cancer cells

DNA in apoptotic cells is sensitive to denaturation by formamide in contrast to necrotic cells. This sensitivity is due to changes in apoptotic cells chromatin. The specific denaturation was detected with monoclonal antibody to single stranded DNA. Using ssDNA Apoptosis

ELISA kit (Chemicon International, Inc.) chimeric compounds were analyzed for their potential to induce apoptosis in PC-3 cells compared to TRAIL (TNF-related apoptosis-inducing ligand), a known apoptotic agent (R&D Systems, Inc.). Results are presented in values of apoptotic index which is calculated by dividing the optical density of treated and untreated cells.

5

Compound (50µM)	Apoptotic index (48 hours)
TRAIL	2.3
6016	3.4
6082	2.6
6086	2.5
6088	1.6
6096	3.2
6102	2.7
6104	2.9
6132	2.0
6134	2.2
6136	2.4
6138	3.6
6140	4.0
6152	1.5
6146	1.8

### CLAIMS

What is claimed is:

- 5           1.     A protein kinase inhibitor comprising a peptide having the sequence

Arg-Pro-Arg-R<sub>4</sub>-R<sub>5</sub>-R<sub>6</sub>-R<sub>7</sub>

wherein:

R<sub>4</sub> is selected from the group consisting of: norvaline, valine, methionine, ornithine, arginine and norleucine;

- 10     R<sub>5</sub> is selected from the group consisting of: tyrosine, threonine, aminobutyric acid and glutamic acid;

R<sub>6</sub> is selected from the group consisting of: diaminopropionic acid, alanine, aminobutyric acid, aspartic acid and diaminobutyric acid; and

R<sub>7</sub> is selected from the group consisting of: homoleucine, phenylalanine and norleucine.

- 15           2.     The protein kinase inhibitor according to claim 1 wherein:

R<sub>4</sub> is norvaline;

R<sub>5</sub> is tyrosine;

R<sub>6</sub> is diaminopropionic acid; and

- 20     R<sub>7</sub> is homoleucine.

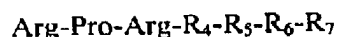
3.     The protein kinase inhibitor according to claim 1 further comprising a moiety W-L connected to a group selected from the group consisting of: the peptide's amino terminus, the peptide's carboxy terminus, a side chain of A<sub>4</sub>, A<sub>5</sub>, R<sub>6</sub> or A<sub>7</sub>; wherein W is an hydrophobic moiety, capable of improving cell permeability; and L is i) a bond selected from the group consisting of: amide, amine, urea, carbamate, hydrazine, sulfonamide; or ii) at least one amino acid selected from the group consisting of: glycine, β-alanine, phenylalanine, aminobutyric acid and aminopentanoic acid.

- 30           4.     The protein kinase inhibitor according to claim 3 wherein W is selected from the group consisting of: substituted or unsubstituted alkylene, aliphatic, aromatic or heterocyclic moiety, of 1-18 carbon atoms; and L is selected from the group consisting of an amide bond, glycine and β-alanine.

5. The compound according to claim 4 wherein W is selected from the group consisting of: myristyl, lauryl, 5-mercaptoaminopropyl-isoquinoline and 5-aminoethylsulfonamide isoquinoline.

- 5 6. The compound according to claim 5 selected from the group consisting of:  
Lauryl-Gly-Arg-Pro-Arg-Nva-Tyr-Dap-Hol;  
Lauryl- $\beta$ -Ala-Arg-Pro-Arg-Nva-Tyr-Dap-Hol;  
Myristyl-Arg-Pro-Arg-Nva-Tyr-Dap-Hol;  
Myristyl-Gly-Arg-Pro-Arg-Nva-Tyr-Dap-Hol;  
10 Myristyl- $\beta$ -Ala-Arg-Pro-Arg-Nva-Tyr-Dap-Hol.

7. A pharmaceutical composition comprising as an active ingredient A protein kinase inhibitor comprising a peptide having the sequence



15 wherein:

R<sub>4</sub> is selected from the group consisting of: norvaline, valine, methionine, ornithine, arginine and norleucine;

R<sub>5</sub> is selected from the group consisting of: tyrosine, threonine, aminobutyric acid and glutamic acid;

20 R<sub>6</sub> is selected from the group consisting of: diaminopropionic acid, alanine, aminobutyric acid, aspartic acid and diaminobutyric acid; and

R<sub>7</sub> is selected from the group consisting of: homoleucine, phenylalanine and norleucine.

8. The pharmaceutical composition according to claim 7 wherein:

25 R<sub>4</sub> is norvaline;

R<sub>5</sub> is tyrosine;

R<sub>6</sub> is diaminopropionic acid;

R<sub>7</sub> is homoleucine; and

a pharmaceutically acceptable diluent or carrier.

30

9. The pharmaceutical composition according to claim 7 wherein the peptide further comprises a moiety W-L connected to a group selected from the group consisting of: the



peptide's amino terminus, the peptide's carboxy terminus, a side chain of A<sub>4</sub>, A<sub>5</sub>, R<sub>6</sub> or A<sub>7</sub>; wherein W is an hydrophobic moiety, capable of improving cell permeability; and L is i) a bond selected from the group consisting of: amide, amine, urea, carbamate, hydrazine, sulfonamide; or ii) at least one amino acid selected from the group consisting of: glycine,  $\beta$ -alanine,

5 phenylalanine, aminobutyric acid and aminopentanoic acid.

10. The pharmaceutical composition according to claim 9 wherein W is selected from the group consisting of: substituted or unsubstituted alkylene, aliphatic, aromatic or heterocyclic moiety, of 1-18 carbon atoms; and L is selected from the group consisting of an amide bond,  
10 glycine and  $\beta$ -alanine.

11. The pharmaceutical composition according to claim 10 wherein W is selected from the group consisting of: myristyl, lauryl, 5-mercaptoaminopropyl-isoquinoline and 5-aminoethylsulfonamide isoquinoline.

15 12. The pharmaceutical composition according to claim 11 wherein the peptide is selected from the group consisting of:

Lauryl-Gly-Arg-Pro-Arg-Nva-Tyr-Dap-Hol;

Lauryl- $\beta$ -Ala-Arg-Pro-Arg-Nva-Tyr-Dap-Hol;

20 Myristyl-Arg-Pro-Arg-Nva-Tyr-Dap-Hol;

Myristyl-Gly-Arg-Pro-Arg-Nva-Tyr-Dap-Hol;

Myristyl- $\beta$ -Ala-Arg-Pro-Arg-Nva-Tyr-Dap-Hol.

13. A method of preparing a medicament for treatment of a disease selected from the  
25 group consisting of: cancer, diabetes, cardiovascular pathology, hemorrhagic shock, obesity, inflammatory disease, disease of the central nervous system, and autoimmune disease, which comprises incorporating a compound according to claim 1 in a pharmaceutical composition, optionally with a pharmaceutically acceptable diluent or excipient.

**PROTEIN KINASE INHIBITORS COMPRISING ATP MIMETICS  
CONJUGATED TO PEPTIDES OR PEPTIDOMIMETICS**

**ABSTRACT**

5        The present invention provides small molecules having high affinity to the ATP binding  
site of protein kinases, which are conjugated to a peptide or peptidomimetic moiety which mimics  
the substrate of PKB. The chimeric compounds according to the present invention preferably  
serve as PKB inhibitors with improved activity and selectivity. Novel ATP mimetic compounds,  
particularly isoquinoline derivatives, conjugated with peptides or peptidomimetics are useful as  
10    inhibitors of protein kinases for experimental, medical, and drug design purposes. Furthermore,  
pharmaceutical compositions comprising these protein kinase inhibitors, and methods of using  
such compositions for treatment and diagnosis of cancers, diabetes, cardiovascular pathologies,  
hemorrhagic shock, obesity, inflammatory diseases, diseases of the central nervous system, and  
autoimmune disease, are disclosed.

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<223> Xaa=Glu-(CO-N-bAla-5-aminoethylsulfoneamide-isoquinoline)

<400> 31

Arg Pro Arg Thr Xaa Ser Phe  
1 5

## 87534-4300 sequence listing.txt

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<222> (4)..(4)  
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<222> (5)..(5)  
<223> Xaa=Glu-(CO-N-bGaba-5-aminoethylsulfoneamide-isoquinoline)

<400> 32

Arg Pro Arg Xaa Xaa Ser Phe  
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<400> 33

Arg Pro Arg Thr Xaa Ser Phe  
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<210> 34  
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<220>  
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<220>  
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<222> (4)..(4)  
<223> Xaa=Nle

<220>  
<221> misc\_feature

## 87534-4300 Sequence Listing.txt

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<223> Xaa=Glu-(CO-N-bAla-5-aminoethylsulfoneamide-isoquinoline)

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<223> Xaa=Nle

<400> 34

Arg Pro Arg Xaa Xaa Ser Xaa  
1 5

<210> 35  
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<220>  
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<400> 35

Arg Pro Arg Thr Xaa Ser Phe  
1 5

<210> 36  
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<220>  
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<220>  
<221> misc\_feature  
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<400> 36

Arg Pro Arg Thr Xaa Ser Phe  
1 5

<210> 37  
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## 87534-4300 sequence listing.txt

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<220>  
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<400> 37

Arg Pro Arg Xaa Xaa Ser Phe  
1 5

<210> 38  
<211> 7  
<212> PRT  
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<220>  
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<220>  
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<220>  
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<400> 39

Arg Pro Arg Xaa Xaa Ser Phe  
1 5

<210> 40  
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## 87534-4300 Sequence listing.txt

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<220>  
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<223> Xaa=Nle

<220>  
<221> misc\_feature  
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<400> 40

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1 5

<210> 41  
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<220>  
<223> synthetic peptide

<220>  
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<223> Xaa=Nle

<220>  
<221> misc\_feature  
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<223> Xaa=Glu-(CO-N-bAla-5-aminoethylsulfoneamide-isoquinoline)

<220>  
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<400> 41

Arg Pro Arg Xaa xaa Xaa Phe  
1 5

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<220>  
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<220>

## 87534-4300 sequence listing.txt

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<223> Xaa=Nle  
  
<220>  
<221> misc\_feature  
<222> (5)..(5)  
<223> Xaa=Glu-(CO-N-bAla-5-aminoethylsulfoneamide-isoquinoline)

<220>  
<221> misc\_feature  
<222> (6)..(6)  
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Arg Pro Arg Xaa Xaa Xaa Phe  
1 5

<210> 43  
<211> 7  
<212> PRT  
<213> Artificial

<220>  
<223> synthetic peptide

<220>  
<221> misc\_feature  
<222> (4)..(4)  
<223> Xaa=Nva

<220>  
<221> misc\_feature  
<222> (5)..(5)  
<223> Xaa=Glu-(CO-N-5-aminoethylsulfoneamide-isoquinoline)

<220>  
<221> misc\_feature  
<222> (7)..(7)  
<223> Xaa=Homoleucine (Hol)

<400> 43

Arg Pro Arg Xaa Xaa Ala Xaa  
1 5

<210> 44  
<211> 7  
<212> PRT  
<213> Artificial

<220>  
<223> synthetic peptide

<220>  
<221> misc\_feature  
<222> (4)..(4)  
<223> Xaa=Nva

87534-4300 sequence listing.txt

<220>  
<221> misc\_feature  
<222> (5)..(5)  
<223> Xaa=Glu-(CO-N-bAla-5-aminoethylsulfoneamide-isoquinoline)

<220>  
<221> misc\_feature  
<222> (7)..(7)  
<223> Xaa=Homoleucine (Ho1)

<400> 44

Arg Pro Arg Xaa Xaa Ala Xaa  
1 5

Figure 1.

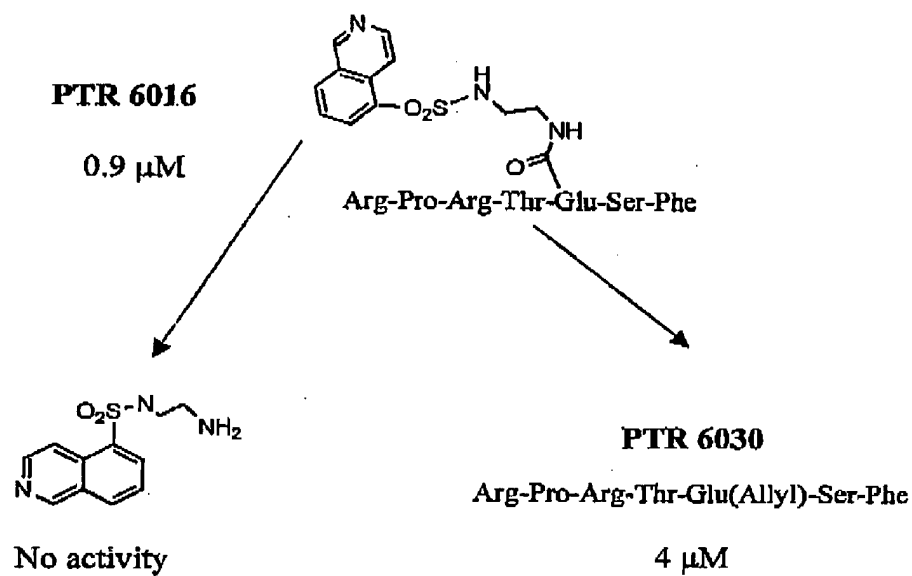


Figure 2.

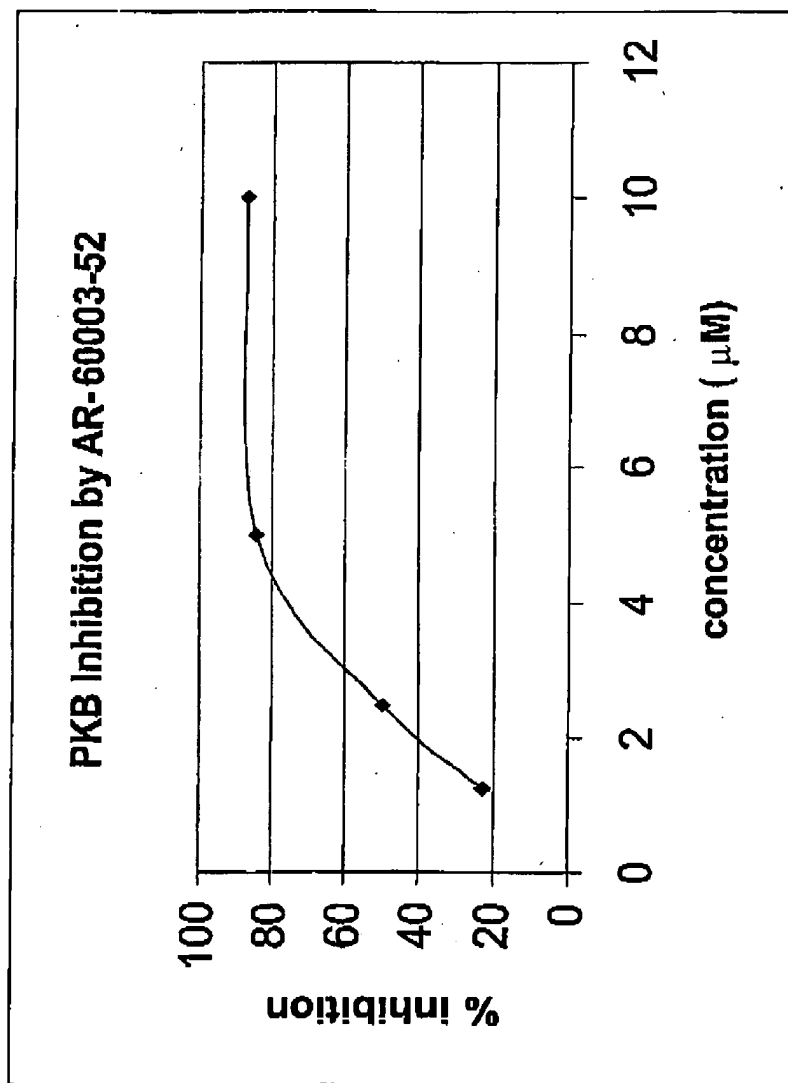


Figure 3

